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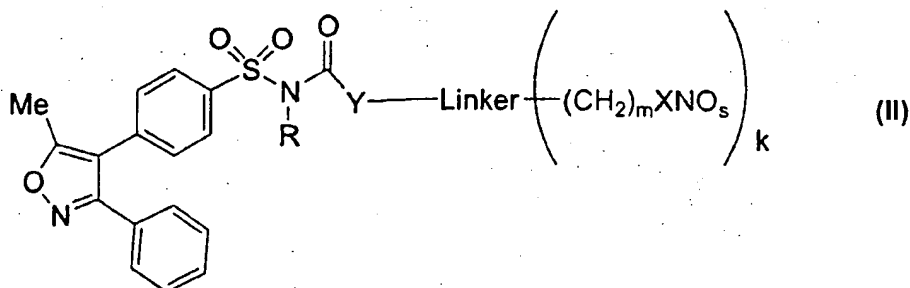
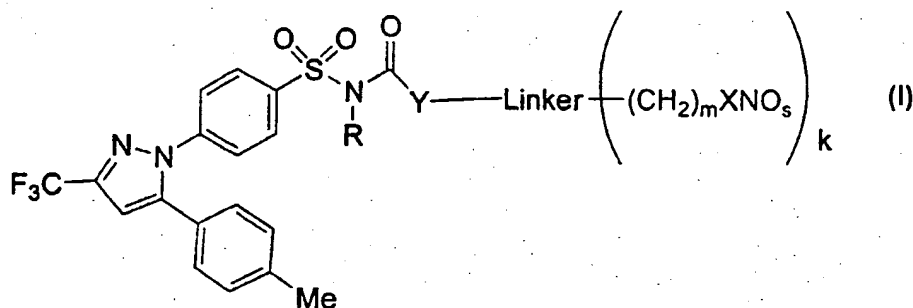
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(54) Title: NITRIC OXIDE RELEASING SELECTIVE CYCLOOXYGENASE-2 INHIBITORS



(57) Abstract: The invention encompasses novel compounds of Formula (I) and Formula (II), which are nitric oxide-releasing prodrugs useful in the treatment of cyclooxygenase-2 mediated diseases. The invention also encompasses certain pharmaceutical compositions and methods for treatment of cyclooxygenase-2 mediated diseases comprising the use of compounds of Formula (I) or Formula (II). The above compounds may be used as a combination therapy with low-dose aspirin to treat chronic cyclooxygenase-2 mediated diseases or conditions while simultaneously reducing the risk of thrombotic cardiovascular events.



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For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

TITLE OF THE INVENTION

NITRIC OXIDE RELEASING SELECTIVE CYCLOOXYGENASE-2 INHIBITORS

BACKGROUND OF THE INVENTION

5 Selective inhibitors of cyclooxygenase-2 are a sub-class of the class of drugs known as non-steroidal antiinflammatory drugs (NSAIDs). The NSAIDs are active in reducing the prostaglandin-induced pain and swelling associated with the inflammation process but are also active in affecting other prostaglandin-regulated processes not associated with the inflammation process. Thus, use of high doses of most common NSAIDs can produce severe side effects, including life threatening ulcers, that limit their therapeutic potential. An
10 alternative to NSAIDs is the use of corticosteroids, which have even more drastic side effects, especially when long term therapy is involved.

 Previous NSAIDs have been found to prevent the production of prostaglandin by inhibiting enzymes in the human arachidonic acid/prostaglandin pathway including the enzyme
15 cyclooxygenase (COX). The discovery that there are two isoforms of the COX enzyme, the first, COX-1, being involved with physiological functions and the second, COX-2, being induced in inflamed tissue, has given rise to a new approach. While conventional NSAIDs block both forms of the enzyme, the identification of the inducible COX-2 enzyme associated with inflammation has provided a viable target of inhibition which more effectively reduces
20 inflammation and produces fewer and less drastic side effects. Many compounds which have activity as COX-2 inhibitors have been identified, including rofecoxib (VIOXX®), etoricoxib (ARCOXIA™), celecoxib (CELEBREX®) and valdecoxib (BEXTRA™), and much research continues in this area.

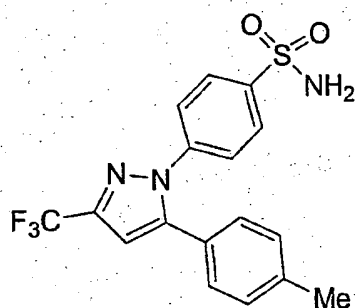
 Many patients with a chronic cyclooxygenase-2 mediated disease or condition are
25 elderly and thus are at increased risk for thrombotic cardiovascular events, such as stroke, myocardial ischemia, myocardial infarction, angina pectoris, transient ischemic attack (TIA; amaurosis fugax), reversible ischemic neurologic deficits, and any similar thrombotic event in any vascular bed (splanchnic, renal, aortic, peripheral, etc.). Moreover, there is evidence that patients with chronic inflammatory conditions, such as rheumatoid arthritis and systemic lupus
30 erythematosus are at increased risk for thrombotic cardiovascular events. Thus, it is desirable that such patients receive appropriate therapy to reduce their risk of such events.

 NO-releasing forms of non-steroidal anti-inflammatory drugs are known in the art and are reported to have improved gastrointestinal and cardiovascular safety profiles over their conventional NSAID counterparts. Furthermore, NO-releasing forms of selective

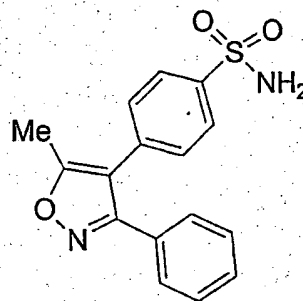
cyclooxygenase-2 selective inhibitors are disclosed in WO 01/45703, published on June 28, 2001, which is hereby incorporated by reference in its entirety.

The present invention provides for novel nitrosated or nitrosylated prodrugs for cyclooxygenase-2 selective inhibitors that are useful for treating cyclooxygenase-2 mediated diseases or conditions which can be administered alone or in combination with low-dose aspirin. Thus, the invention provides for a clearly superior profile than that hitherto obtainable in that it provides efficacy in treating chronic cyclooxygenase-2 mediated diseases or conditions, effectively reducing the risk of thrombotic cardiovascular events and renal side effects and at the same time reduces the risk of GI ulceration or bleeding.

In particular, the present invention provides novel prodrugs of celebrex (Compound A) and Valdecoxib (Compound B):



Compound A



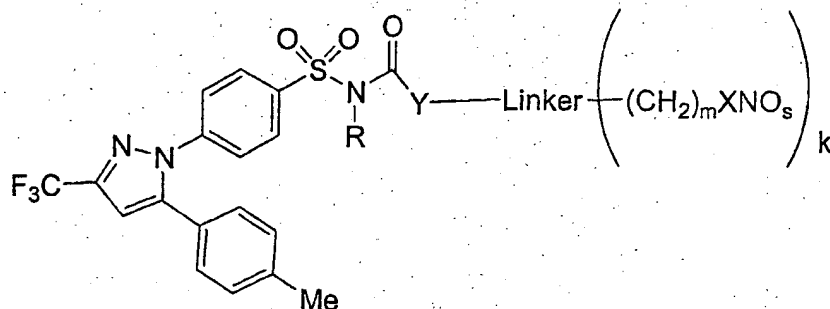
Compound B

Celebrex, methods of preparing celecoxib and methods of using celecoxib are disclosed in U.S. Patent 5,466,823, issued November 14, 1995, which is hereby incorporated by reference.

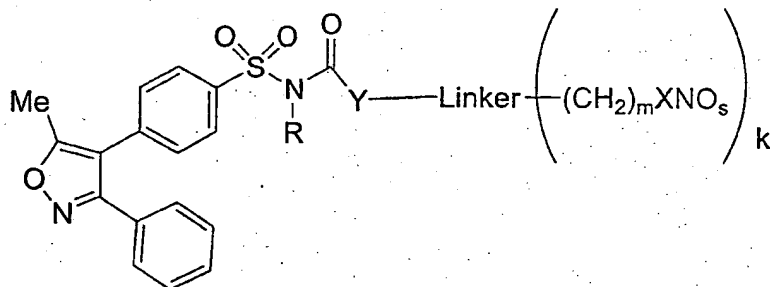
Valdecoxib, methods of preparing valdecoxib and methods of using valdecoxib are disclosed in U.S. Patent U.S. 5,633,272, issued May 27, 1997, which is hereby incorporated by reference.

SUMMARY OF THE INVENTION

The invention encompasses novel compounds of Formula I and Formula II, which are nitric oxide-releasing prodrugs useful in the treatment of cyclooxygenase-2 mediated diseases.



Formula I

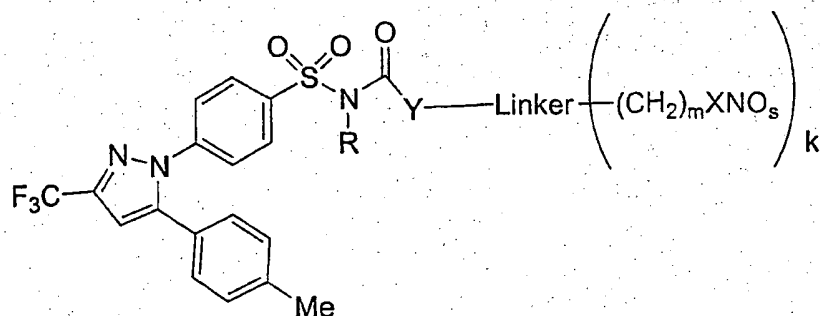


Formula II

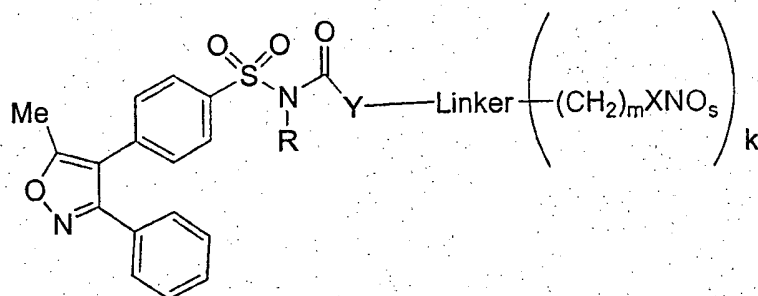
The invention also encompasses certain pharmaceutical compositions and methods for treatment of cyclooxygenase-2 mediated diseases comprising the use of compounds of Formula I or Formula II. The above compounds may be used as a combination therapy with low-dose aspirin to treat chronic cyclooxygenase-2 mediated diseases or conditions while simultaneously reducing the risk of thrombotic cardiovascular events.

DETAILED DESCRIPTION OF THE INVENTION

The invention encompasses novel compounds of Formula I and Formula II, which are nitric oxide-releasing prodrugs useful in the treatment of cyclooxygenase-2 mediated diseases.



Formula I



Formula II

or a pharmaceutically acceptable salt thereof wherein

each s is independently 1 or 2;

k is 1, 2, 3 or 4;

each m is independently 0, 1, 2, 3 or 4;

each X is independently O or S;

Y is a bond, S, O or NR_1 , wherein R_1 is hydrogen or C_{1-6} alkyl;

R is hydrogen or C_{1-6} alkyl;

the Linker is selected from the group consisting of:

(a) $-(CH_2)_n$, wherein n is 0, 1, 2, 3 or 4,

(b) C_{3-6} cycloalkyl, wherein the C_{3-6} cycloalkyl optionally mono-, di- or tri-substituted with a substituent selected from the group consisting of (1) halo, (2) C_{1-3} alkyl, (3) C_{1-3} alkoxy,

(4) Hydroxy,

(5) NO₂,

(6) CO₂,

(7) CF₃,

(8) CN;

(9) CH₂COOH

(10) CH₂COO-C₁₋₃alkyl,

(11) C₁₋₃alkthio,

(c) aryl, wherein the aryl is selected from the group consisting of phenyl and naphthyl, wherein the aryl is optionally mono-, di- or tri-substituted with a substituent selected from the group consisting of

(1) halo,

(2) C₁₋₃alkyl,

(3) C₁₋₃alkoxy,

(4) Hydroxy,

(5) NO₂,

(6) CO₂,

(7) CF₃,

(8) CN;

(9) CH₂COOH

(10) CH₂COO-C₁₋₃alkyl,

(11) C₁₋₃alkthio,

(d) Heteroaryl optionally mono-, di- or tri- substituted with substituents selected from the group consisting of,

(1) halo,

(2) C₁₋₃alkyl,

(3) C₁₋₃alkoxy,

(4) Hydroxy,

(5) NO₂,

(6) CO₂,

(7) CF₃,

(8) CN;

(9) CH₂COOH

(10) CH₂COO-C₁₋₃alkyl,

(11) C₁₋₃alkthio.

For purposes of this specification, heteroaryl or benzoheteroaryl group includes benzimidazolyl, benzofuranyl, benzopyrazolyl, benzotriazolyl, benzothiophenyl, benzoxazolyl, carbazolyl, carbolinyl, cinnolinyl, furanyl, imidazolyl, indolinyl, indolyl, indolazinyl, indazolyl, isobenzofuranyl, isoindolyl, isoquinolyl, isothiazolyl, isoxazolyl, naphthyridinyl, oxadiazolyl, oxazolyl, pyrazinyl, pyrazolyl, pyridopyridinyl, pyridazinyl, pyridyl, pyrimidyl, pyrrolyl, quinazolinyl, quinolyl, quinoxalinyl, thiadiazolyl, thiazolyl, thienyl, triazolyl, azetidyl, 1,4-dioxanyl, hexahydroazepinyl, piperazinyl, piperidinyl, pyrrolidinyl, morpholinyl, thiomorpholinyl, dihydrobenzimidazolyl, dihydrobenzofuranyl, dihydrobenzothiophenyl, dihydrobenzoxazolyl, dihydrofuranyl, dihydroimidazolyl, dihydroindolyl, dihydroisooxazolyl, dihydroisothiazolyl, dihydrooxadiazolyl, dihydrooxazolyl, dihydropyrazinyl, dihydropyrazolyl, dihydropyridinyl, dihydropyrimidinyl, dihydropyrrolyl, dihydroquinolinyl, dihydrotetrazolyl, dihydrothiadiazolyl, dihydrothiazolyl, dihydrothienyl, dihydrotriazolyl, dihydroazetidyl, methylenedioxybenzoyl, tetrahydrofuranyl, and tetrahydrothienyl.

The present compounds are nitric oxide releasing prodrugs which liberate nitric oxide and celecoxib or valdecoxib *in vivo* and can be administered alone or in combination with low dose aspirin. Thus, the invention provides for a clearly superior profile than that hitherto obtainable in that it provides efficacy in treating chronic cyclooxygenase-2 mediated diseases or conditions, effectively reducing the risk of thrombotic cardiovascular events and renal side effects and at the same time reduces the risk of GI ulceration or bleeding.

An embodiment of the invention encompasses a compound of Formula I and Formula II wherein -S(O)₂NH₂ is replaced with S(O)₂CH₃.

Within this embodiment of the invention is encompassed a compound of Formula I wherein:

s is 2;

k is 1;

m is 1 or 2.

Also this embodiment of the invention is encompassed a compound of Formula I wherein R is hydrogen.

Also this embodiment of the invention is encompassed a compound of Formula I wherein X is O.

Also this embodiment of the invention is encompassed a compound of Formula I wherein Y is a bond.

5 Also this embodiment of the invention is encompassed a compound of Formula I wherein k is 1.

Also this embodiment of the invention is encompassed a compound of Formula I wherein m is 1.

10 Also this embodiment of the invention is encompassed a compound of Formula I wherein
R is hydrogen;
Y is a bond;
s is 2;
k is 1; and
15 m is 1.

Also within this embodiment of the invention is encompassed a compound of Formula I wherein the Linker is $-(CH_2)_n$, wherein n is 1 or 2.

20 Also within this embodiment of the invention is encompassed a compound of Formula I wherein the Linker is C₃-6cycloalkyl, wherein the C₃-6cycloalkyl optionally mono-, di- or tri-substituted with a substituent selected from the group consisting of

- 25 (1) halo,
(2) Methyl,
(3) Methoxy,
(4) Hydroxy,
(5) NO₂,
(6) CO₂,
30 (7) CF₃,
(8) CN;
(9) CH₂COOH

Also within this embodiment of the invention is encompassed a compound of Formula I wherein the Linker is aryl, wherein the aryl is selected from the group consisting of phenyl and naphthyl, wherein the aryl is optionally mono-, di- or tri-substituted with a substituent selected from the group consisting of

- (1) halo,
- (2) Methyl,
- (3) Methoxy,
- (4) Hydroxy,
- (5) NO₂,
- (6) CO₂,
- (7) CF₃,
- (8) CN,
- (9) CH₂COOH.

Also within this embodiment of the invention is encompassed a compound of Formula I

and Formula II wherein the Linker is benzimidazolyl, benzofuranyl, benzopyrazolyl, benzotriazolyl, benzothiophenyl, benzoxazolyl, carbazolyl, carbolinyl, cinnolinyl, furanyl, imidazolyl, indolinyl, indolyl, indolazinyl, indazolyl, isobenzofuranyl, isoindolyl, isoquinolyl, isothiazolyl, isoxazolyl, naphthyridinyl, oxadiazolyl, oxazolyl, pyrazinyl, pyrazolyl, pyridopyridinyl, pyridazinyl, pyridyl, pyrimidyl, pyrrolyl, quinazolinyl, quinolyl, quinoxalinyl, thiadiazolyl, thiazolyl, thienyl, triazolyl, azetidyl, 1,4-dioxanyl, hexahydroazepinyl, piperazinyl, piperidinyl, pyrrolidinyl, morpholinyl, thiomorpholinyl, dihydrobenzimidazolyl, dihydrobenzofuranyl, dihydrobenzothiophenyl, dihydrobenzoxazolyl, dihydrofuranyl, dihydroimidazolyl, dihydroindolyl, dihydroisooxazolyl, dihydroisothiazolyl, dihydrooxadiazolyl, dihydrooxazolyl, dihydropyrazinyl, dihydropyrazolyl, dihydropyridinyl, dihydropyrimidinyl, dihydropyrrolyl, dihydroquinolinyl, dihydrotetrazolyl, dihydrothiadiazolyl, dihydrothiazolyl, dihydrothienyl, dihydrotriazolyl, dihydroazetidyl, methylenedioxybenzoyl, tetrahydrofuranyl, and tetrahydrothienyl, optionally mono-, di- or tri- substituted with substituents selected from the group consisting of,

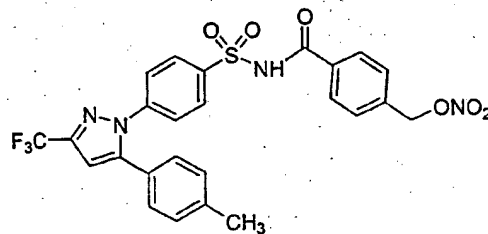
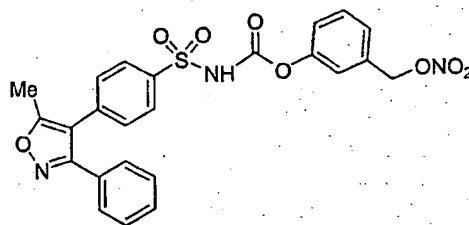
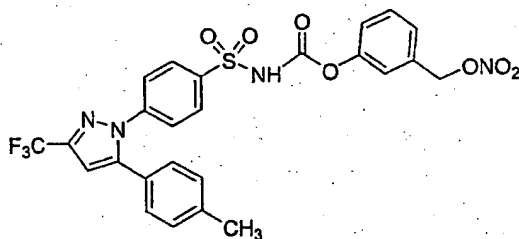
- (1) halo,
- (2) Methyl,
- (3) Methoxy,

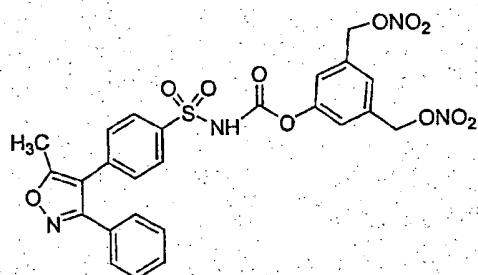
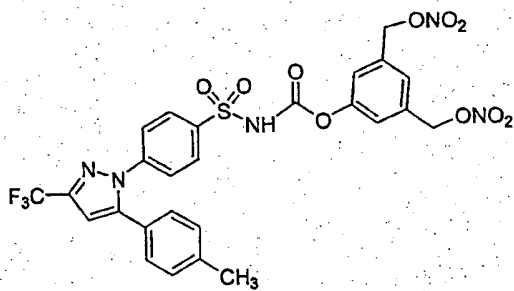
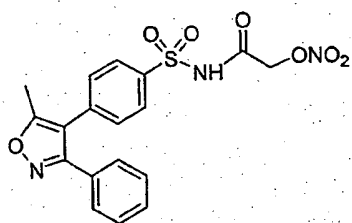
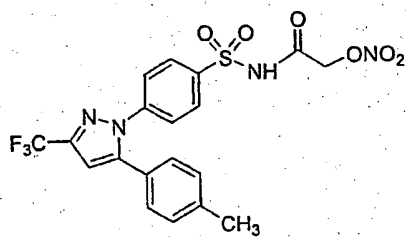
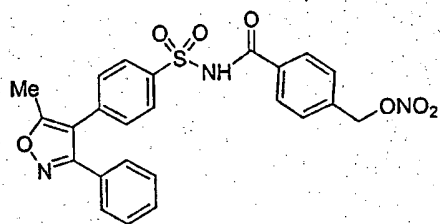
- (4) Hydroxy,
(5) NO₂,
(6) CO₂,
(7) CF₃,
(8) CN;
(9) CH₂COOH

Another embodiment of the invention encompasses the compound of Formula I or II wherein the Linker is pyridyl optionally substituted as above.

Another embodiment of the invention encompasses a compound of Formula I or Formula II wherein s is 2.

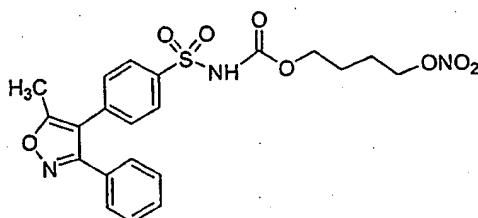
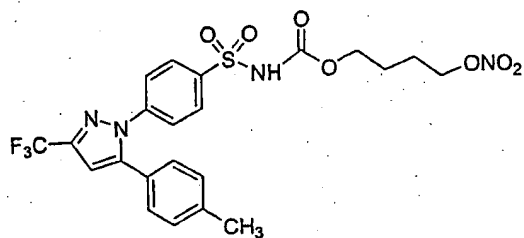
Another embodiment of the invention encompasses a compound selected from the following group:





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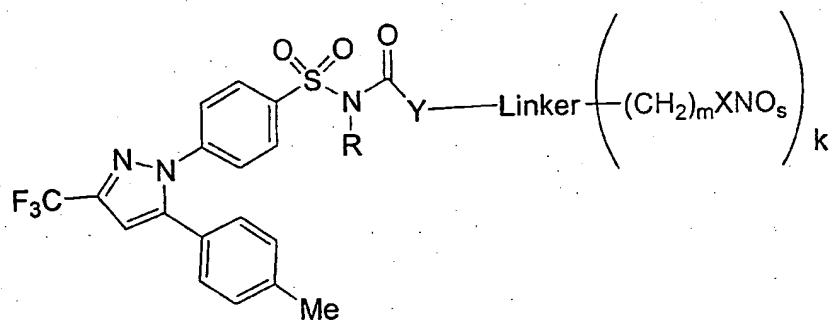
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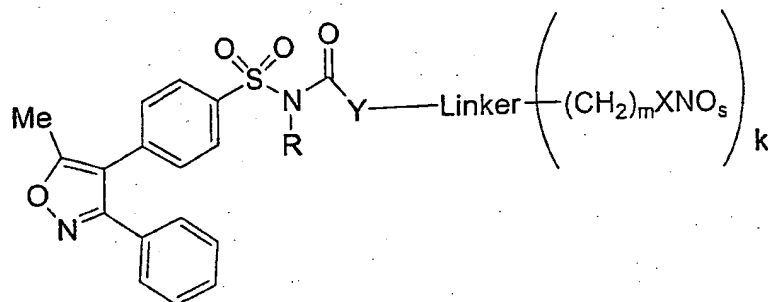
Another embodiment of the invention encompasses a compound of Formula I or

Formula II



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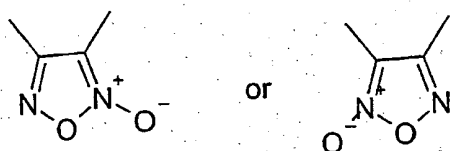
Formula I



Formula II

or a pharmaceutically acceptable salt thereof wherein

- 5 each s is independently 1 or 2;
 k is 1, 2, 3 or 4;
 each m is independently 0, 1, 2, 3 or 4;
 each X is independently O or S;
 Y is a bond, S, O or NR₁, wherein R₁ is hydrogen or C₁-6alkyl;
 10 R is hydrogen or C₁-6alkyl;
 the Linker is selected from the group consisting of:



- 15 The invention also encompasses a pharmaceutical composition comprising a compound of Formula I or Formula II and a pharmaceutically acceptable carrier.

- The invention also encompasses a method of treating an inflammatory disease susceptible to treatment with a non-steroidal anti-inflammatory agent comprising administering to a patient in need of such treatment of a non-toxic therapeutically effective amount of a compound of Formula I or Formula II. Within this embodiment is encompassed the above method wherein the patient is also at risk of a thrombotic cardiovascular event.
- 20

- Another embodiment of the invention encompasses method of treating cyclooxygenase mediated diseases advantageously treated by an active agent that selectively inhibits COX-2 in preference to COX-1 comprising administering to a patient in need of such treatment of a non-toxic therapeutically effective amount of a compound of Formula I. Within this embodiment is encompassed the above method wherein the patient is also at risk of a thrombotic cardiovascular event.
- 25

- Another embodiment of the invention encompasses a method for treating a chronic cyclooxygenase-2 mediated disease or condition and reducing the risk of a thrombotic cardiovascular event in a human patient in need of such treatment and at risk of a thrombotic cardiovascular event comprising orally concomitantly or sequentially administering to said
- 30

patient a compound of Formula I in an amount effective to treat the cyclooxygenase-2 mediated disease or condition and aspirin in an amount effective to reduce the risk of the thrombotic cardiovascular event. Within this embodiment is encompassed the above method wherein the compound of Formula I is administered orally on a once daily basis. Within this embodiment is encompassed the above method wherein the compound of Formula I or Formula II is administered orally on a twice daily basis. Within this embodiment is encompassed the above method wherein the cyclooxygenase-2 selective mediated disease or condition is selected from the group consisting of: osteoarthritis, rheumatoid arthritis and chronic pain. Within this embodiment is encompassed the above method wherein aspirin is administered at a dose of about 30 mg to about 1 g. Within this embodiment is encompassed the above method wherein aspirin is administered at a dose of about 80 to about 650 mg. Within this embodiment is encompassed the above method wherein aspirin is administered at a dose of about 81 mg or about 325 mg. Within this embodiment is encompassed the above method wherein aspirin is orally administered once daily.

The invention also encompasses a pharmaceutical composition comprising a compound of Formula I or Formula II and aspirin in combination with a pharmaceutically acceptable carrier.

For purposes of this specification alkyl is defined to include linear, branched, and cyclic structures, with C₁₋₆alkyl including including methyl, ethyl, propyl, 2-propyl, s- and t-butyl, butyl, pentyl, hexyl, 1,1-dimethylethyl, cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl. Similarly, C₁₋₆alkoxy is intended to include alkoxy groups of from 1 to 6 carbon atoms of a straight, branched, or cyclic configuration. Examples of lower alkoxy groups include methoxy, ethoxy, propoxy, isopropoxy, cyclopropyloxy, cyclohexyloxy, and the like. Likewise, C₁₋₆alkylthio is intended to include alkylthio groups of from 1 to 6 carbon atoms of a straight, branched or cyclic configuration. Examples of lower alkylthio groups include methylthio, propylthio, isopropylthio, cycloheptylthio, etc. By way of illustration, the propylthio group signifies -SCH₂CH₂CH₃.

Some of the compounds described herein contain olefinic double bonds, and unless specified otherwise, are meant to include both E and Z geometric isomers.

The term "treating a chronic cyclooxygenase-2 mediated disease or condition" means treating or preventing any chronic disease or condition that is advantageously treated or prevented by inhibiting the cyclooxygenase-2 enzyme. The term includes the relief of pain, fever and inflammation of a variety of conditions including rheumatic fever, symptoms associated with influenza or other viral infections, common cold, low back pain, neck pain,

dysmenorrhea, headache, migraine, toothache, sprains and strains, myositis, neuralgia, synovitis, arthritis, including rheumatoid arthritis, degenerative joint diseases (osteoarthritis), gout, ankylosing spondylitis, bursitis, burns, injuries, and pain and inflammation following surgical procedures. In addition, such a compound may inhibit cellular neoplastic transformations and metastatic tumor growth and hence can be used in the treatment and/or prevention of cancer. In addition, such a compound may inhibit the onset or progression of Alzheimer's disease or cognitive impairment. The term also includes the treatment and/or prevention of cyclooxygenase-mediated proliferative disorders such as may occur in diabetic retinopathy and tumor angiogenesis. The term "treating" encompasses not only treating a patient to relieve the patient of the signs and symptoms of the disease or condition but also prophylactically treating an asymptomatic patient to prevent the onset or progression of the disease or condition.

A "thrombotic cardiovascular event" is defined as any sudden event of a type known to be caused by platelet aggregation, thrombosis, and subsequent ischemic clinical events, including thrombotic or thromboembolic stroke, myocardial ischemia, myocardial infarction, angina pectoris, transient ischemic attack (TIA; amaurosis fugax), reversible ischemic neurologic deficits, and any similar thrombotic event in any vascular bed (splanchnic, renal, aortic, peripheral, etc.).

The term "patient in need of such treatment and at risk of a thrombotic cardiovascular event" means a patient in need of both treatment for a cyclooxygenase-2 mediated disease and also at risk of a thrombotic cardiovascular event. One skilled in the art can diagnose a patient that is in need of treatment for a cyclooxygenase-2 mediated disease or condition and also at risk of suffering a thrombotic cardiovascular event. For example, such a patient may be over the age of 50 with osteoarthritis and with a previous myocardial infarction. Other risk factors for a thrombotic cardiovascular event include hypertension, hypercholesterolemia, diabetes mellitus, chronic renal impairment, smoking, and any prior personal or family history of such an event. Administration of the drug combination to the patient includes both self-administration and administration to the patient by another person.

The terms "nitric oxide releasing-cyclooxygenase-2 selective inhibitor," "NO-cyclooxygenase-2 selective inhibitor," "nitric oxide releasing-COX-2 inhibitor" and "NO-COX-2 inhibitor" mean a modified version of a cyclooxygenase-2 selective inhibitor or a prodrug as defined above linked to a NO releasing moiety by means of a linking group such as an ester linkage.

The term "amounts that are effective to treat" is intended to mean that amount of a drug or pharmaceutical agent that will elicit the biological or medical response of a tissue, a

system, animal or human that is being sought by a researcher, veterinarian, medical doctor or other clinician. The term also encompasses the amount of a pharmaceutical drug that will prevent or reduce the risk of occurrence of the biological or medical event that is sought to be prevented in a tissue, a system, animal or human by a researcher, veterinarian, medical doctor or other clinician. The inhibitor of cyclooxygenase-2 may be administered at a dosage level up to conventional dosage levels for NSAIDs. Suitable dosage levels will depend upon the antiinflammatory effect of the chosen inhibitor of cyclooxygenase-2, but typically suitable levels will be about 0.01 to about 50 mg/kg per day. The compound may be administered on a regimen of once or twice per day.

The term "amount effective to reduce the risk of" means the amount of a pharmaceutical drug that will prevent or reduce the risk of occurrence of the biological or medical event that is sought to be prevented in a tissue, a system, animal or human by a researcher, veterinarian, medical doctor or other clinician. Aspirin is administered at a dose of about 30 mg to about 1 g once daily, preferably at a dose of about 80 mg to about 650 mg.

The term "concomitantly administering" means administering the agents substantially concurrently. The term "concomitantly administering" encompasses not only administering the two agents in a single pharmaceutical dosage form but also the administration of each active agent in its own separate pharmaceutical dosage formulation. Where separate dosage formulations are used, the agents can be administered at essentially the same time, i.e., concurrently.

The term "sequentially administering" means administering the agents at separately staggered times. Thus, agents can be sequentially administered such that the beneficial pharmaceutical effect of NO-aspirin and the COX-2 inhibitor or aspirin and the NO-COX-2 inhibitor are realized by the patient at substantially the same time. Thus, for example, if a COX-2 selective inhibitor and NO releasing aspirin are both administered on a once a day basis, the interval of separation between sequential administration of the two agents can be up to twelve hours apart.

The pharmaceutical compositions of the present invention comprise a compound of Formula I as an active ingredient or a pharmaceutically acceptable salt, thereof, and may also contain a pharmaceutically acceptable carrier and optionally other therapeutic ingredients. The term "pharmaceutically acceptable salts" refers to salts prepared from pharmaceutically acceptable non-toxic bases including inorganic bases and organic bases. Salts derived from inorganic bases include aluminum, ammonium, calcium, copper, ferric, ferrous, lithium, magnesium, manganic salts, manganous, potassium, sodium, zinc, and the like. Particularly

preferred are the ammonium, calcium, magnesium, potassium, and sodium salts. Salts derived from pharmaceutically acceptable organic non-toxic bases include salts of primary, secondary, and tertiary amines, substituted amines including naturally occurring substituted amines, cyclic amines, and basic ion exchange resins, such as arginine, betaine, caffeine, choline, N,N-dibenzylethylenediamine, diethylamine, 2-diethylaminoethanol, 2-dimethylaminoethanol, ethanolamine, ethylenediamine, N-ethylmorpholine, N-ethylpiperidine, glucamine, glucosamine, histidine, hydrabamine, isopropylamine, lysine, methylglucamine, morpholine, piperazine, piperidine, polyamine resins, procaine, purines, theobromine, triethylamine, trimethylamine, tripropylamine, tromethamine, and the like.

It will be understood that in the discussion of methods of treatment which follows, references to the compounds of Formula I or Formula II are meant to also include the pharmaceutically acceptable salts.

The Compound of Formula I and Formula II is useful for the relief of pain, fever and inflammation of a variety of conditions including rheumatic fever, symptoms associated with influenza or other viral infections, common cold, low back and neck pain, dysmenorrhea, headache, toothache, sprains and strains, myositis, neuralgia, synovitis, arthritis, including rheumatoid arthritis degenerative joint diseases (osteoarthritis), gout and ankylosing spondylitis, bursitis, burns, injuries, following surgical and dental procedures. In addition, such a compound may inhibit cellular neoplastic transformations and metastatic tumor growth and hence can be used in the treatment of cancer. Compounds of Formula I may also be useful for the treatment of dementia including pre-senile and senile dementia, and in particular, dementia associated with Alzheimer Disease (i.e. Alzheimer's dementia).

Compounds of Formula I and Formula II will also inhibit prostanoid-induced smooth muscle contraction by preventing the synthesis of contractile prostanoids and hence may be of use in the treatment of dysmenorrhea, premature labor and asthma. They will also be useful to inhibit bone loss (osteoporosis).

By virtue of its high cyclooxygenase-2 (COX-2) activity and/or its selectivity for cyclooxygenase-2 over cyclooxygenase-1 (COX-1) as defined above, compounds of Formula I and Formula II will prove useful as an alternative to conventional non-steroidal antiinflammatory drugs (NSAID'S) particularly where such non-steroidal antiinflammatory drugs may be contra-indicated such as in patients with peptic ulcers, gastritis, regional enteritis, ulcerative colitis, diverticulitis or with a recurrent history of gastrointestinal lesions; GI bleeding, coagulation disorders including anemia such as hypoprothrombinemia, haemophilia or other bleeding problems (including those relating to reduced or impaired platelet function);

kidney disease (e.g. impaired renal function); those prior to surgery or taking anticoagulants; and those susceptible to NSAID induced asthma.

Similarly, compounds of Formula I and Formula II, will be useful as a partial or complete substitute for conventional NSAID'S in preparations wherein they are presently co-administered with other agents or ingredients. Thus in further aspects, the invention encompasses pharmaceutical compositions for treating cyclooxygenase-2 mediated diseases as defined above comprising a non-toxic therapeutically effective amount of the compound of Formula I or Formula II as defined above and one or more ingredients such as another pain reliever including acetaminophen or phenacetin; a potentiator including caffeine; an H2-antagonist, aluminum or magnesium hydroxide, simethicone, a decongestant including phenylephrine, phenylpropanolamine, pseudophedrine, oxymetazoline, ephinephrine, naphazoline, xylometazoline, propylhexedrine, or levo-desoxyephedrine; an antiitussive including codeine, hydrocodone, caramiphen, carbetapentane, or dexamethorphan; a diuretic; a sedating or non-sedating antihistamine. In addition the invention encompasses a method of treating cyclooxygenase mediated diseases comprising: administration to a patient in need of such treatment a non-toxic therapeutically effect amount of the compound of Formula I or Formula II, optionally co-administered with one or more of such ingredients as listed immediately above.

Compounds of the present invention are inhibitors of cyclooxygenase-2 and are thereby useful in the treatment of cyclooxygenase-2 mediated diseases as enumerated above. This activity is illustrated by their ability to selectively inhibit cyclooxygenase-2 over cyclooxygenase-1. Accordingly, in one assay, the ability of the compounds of this invention to treat cyclooxygenase mediated diseases can be demonstrated by measuring the amount of prostaglandin E2 (PGE2) synthesized in the presence of arachidonic acid, cyclooxygenase-1 or cyclooxygenase-2 and a compound of Formula I. The IC50 values represent the concentration of inhibitor required to return PGE2 synthesis to 50% of that obtained as compared to the uninhibited control. For the treatment of any of these cyclooxygenase mediated diseases, compounds of Formula I and Formula II may be administered orally, topically, parenterally, by inhalation spray or rectally in dosage unit formulations containing conventional non-toxic pharmaceutically acceptable carriers, adjuvants and vehicles. The term parenteral as used herein includes subcutaneous injections, intravenous, intramuscular, intrasternal injection or infusion techniques. In addition to the treatment of warm-blooded animals such as mice, rats, horses, cattle sheep, dogs, cats, etc., the compound of the invention is effective in the treatment of humans.

As indicated above, pharmaceutical compositions for treating cyclooxygenase-2 mediated diseases as defined may optionally include one or more ingredients as listed above.

The pharmaceutical compositions containing the active ingredient may be in a form suitable for oral use, for example, as tablets, troches, lozenges, aqueous or oily suspensions, dispersible powders or granules, emulsions, hard or soft capsules, or syrups or elixirs. Compositions intended for oral use may be prepared according to any method known to the art for the manufacture of pharmaceutical compositions and such compositions may contain one or more agents selected from the group consisting of sweetening agents, flavoring agents, coloring agents and preserving agents in order to provide pharmaceutically elegant and palatable preparations. Tablets contain the active ingredient in admixture with non-toxic pharmaceutically acceptable excipients which are suitable for the manufacture of tablets. These excipients may be for example, inert diluents, such as calcium carbonate, sodium carbonate, lactose, calcium phosphate or sodium phosphate; granulating and disintegrating agents, for example, corn starch, or alginic acid; binding agents, for example starch, gelatin or acacia, and lubricating agents, for example, magnesium stearate, stearic acid or talc. The tablets may be uncoated or they may be coated by known techniques to delay disintegration and absorption in the gastrointestinal tract and thereby provide a sustained action over a longer period. For example, a time delay material such as glyceryl monostearate or glyceryl distearate may be employed. They may also be coated by the technique described in the U.S. Patent 4,256,108; 4,166,452; and 4,265,874 to form osmotic therapeutic tablets for control release.

Formulations for oral use may also be presented as hard gelatin capsules wherein the active ingredient is mixed with an inert solid diluent, for example, calcium carbonate, calcium phosphate or kaolin, or as soft gelatin capsules wherein the active ingredients is mixed with water or an oil medium, for example peanut oil, liquid paraffin, or olive oil.

Aqueous suspensions contain the active material in admixture with excipients suitable for the manufacture of aqueous suspensions. Such excipients are suspending agents, for example sodium carboxymethyl-cellulose, methylcellulose, hydroxypropylmethy-cellulose, sodium alginate, polyvinyl-pyrrolidone, gum tragacanth and gum acacia; dispersing or wetting agents may be a naturally-occurring phosphatide, for example lecithin, or condensation products of an alkylene oxide with fatty acids, for example polyoxyethylene stearate, or condensation products of ethylene oxide with long chain aliphatic alcohols, for example heptadecaethylene-oxyctanol, or condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as polyoxyethylene sorbitol monooleate, or condensation products of ethylene oxide with partial esters derived from fatty acids and hexitol anhydrides, for example

polyethylene sorbitan monooleate. The aqueous suspensions may also contain one or more preservatives, for example ethyl, or n-propyl, p-hydroxybenzoate, one or more coloring agents, one or more flavoring agents, and one or more sweetening agents, such as sucrose, saccharin or aspartame.

5 Oily suspensions may be formulated by suspending the active ingredient in a vegetable oil, for example arachis oil, olive oil, sesame oil or coconut oil, or in mineral oil such as liquid paraffin. The oily suspensions may contain a thickening agent, for example beeswax, hard paraffin or cetyl alcohol. Sweetening agents such as those set forth above, and flavoring agents may be added to provide a palatable oral preparation. These compositions may be
10 preserved by the addition of an anti-oxidant such as ascorbic acid.

Dispersible powders and granules suitable for preparation of an aqueous suspension by the addition of water provide the active ingredient in admixture with a dispersing or wetting agent, suspending agent and one or more preservatives. Suitable dispersing or wetting agents and suspending agents are exemplified by those already mentioned above.
15 Additional excipients, for example sweetening, flavoring and coloring agents, may also be present.

The pharmaceutical compositions of the invention may also be in the form of an oil-in-water emulsions. The oily phase may be a vegetable oil, for example olive oil or arachis oil, or a mineral oil, for example liquid paraffin or mixtures of these. Suitable emulsifying
20 agents may be naturally-occurring phosphatides, for example soy bean, lecithin, and esters or partial esters derived from fatty acids and hexitol anhydrides, for example sorbitan monooleate, and condensation products of the said partial esters with ethylene oxide, for example polyoxyethylene sorbitan monooleate. The emulsions may also contain sweetening and flavouring agents.

25 Syrups and elixirs may be formulated with sweetening agents, for example glycerol, propylene glycol, sorbitol or sucrose. Such formulations may also contain a demulcent, a preservative and flavoring and coloring agents. The pharmaceutical compositions may be in the form of a sterile injectable aqueous or oleagenous suspension. This suspension may be formulated according to the known art using those suitable dispersing or wetting agents
30 and suspending agents which have been mentioned above. The sterile injectable preparation may also be a sterile injectable solution or suspension in a non-toxic parenterally-acceptable diluent or solvent, for example as a solution in 1,3-butane diol. Among the acceptable vehicles and solvents that may be employed are water, Ringer's solution and isotonic sodium chloride solution. In addition, sterile, fixed oils are conventionally employed as a solvent or suspending

medium. For this purpose any bland fixed oil may be employed including synthetic mono- or diglycerides. In addition, fatty acids such as oleic acid find use in the preparation of injectables.

Compounds of Formula I may also be administered in the form of suppositories for rectal administration of the drug. These compositions can be prepared by mixing the drug with a suitable non-irritating excipient which is solid at ordinary temperatures but liquid at the rectal temperature and will therefore melt in the rectum to release the drug. Such materials are cocoa butter and polyethylene glycols.

For topical use, creams, ointments, jellies, solutions or suspensions, etc., containing the compound of Formula I or Formula II are employed. (For purposes of this application, topical application shall include mouth washes and gargles.)

Dosage levels of the order of from about 0.01 mg to about 50 mg/kg of body weight per day are useful in the treatment of the above-indicated conditions, or alternatively about 0.5 mg to about 2 g per patient per day. For example, inflammation may be effectively treated by the administration of from about 0.01 to 50 mg of the compound per kilogram of body weight per day, or alternatively about 0.5 mg to about 2 g per patient per day.

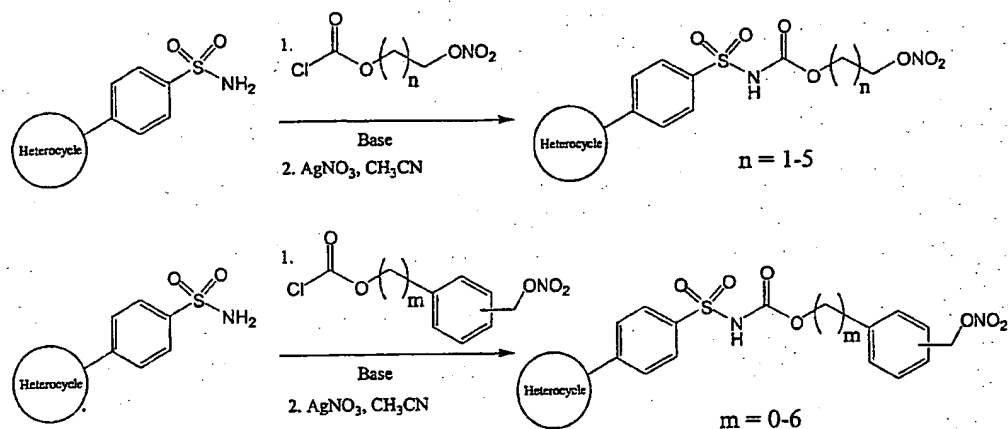
The amount of active ingredient that may be combined with the carrier materials to produce a single dosage form will vary depending upon the host treated and the particular mode of administration. For example, a formulation intended for the oral administration of humans may contain from 0.5 mg to 2 g of active agent compounded with an appropriate and convenient amount of carrier material which may vary from about 5 to about 95 percent of the total composition. Dosage unit forms will generally contain between from about 1 mg to about 500 mg of an active ingredient, typically 25 mg, 50 mg, 100 mg, 200 mg, 300 mg, 400 mg, 500 mg, 600 mg, 800 mg, or 1000 mg.

It will be understood, however, that the specific dose level for any particular patient will depend upon a variety of factors including the age, body weight, general health, sex, diet, time of administration, route of administration, rate of excretion, drug combination and the severity of the particular disease undergoing therapy.

Methods of Synthesis

Method A

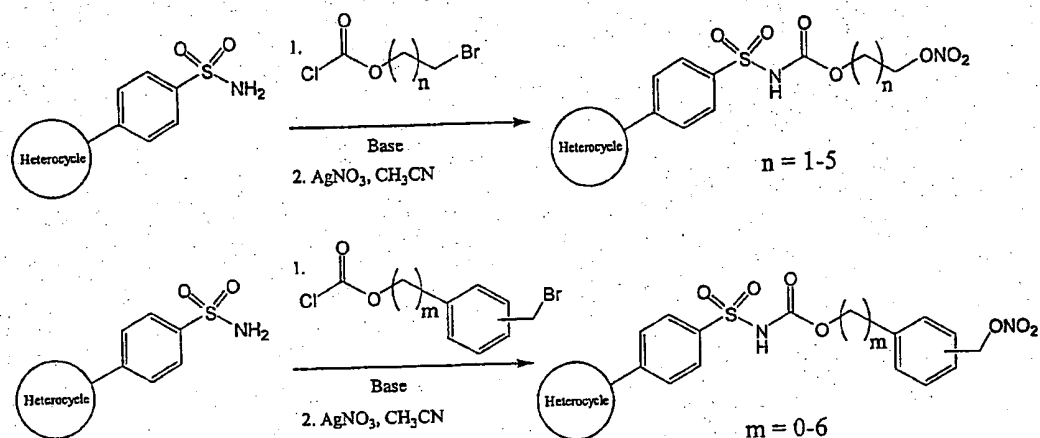
5 Scheme 1



The sulfonylcarbamate derivatives can be prepared according to Scheme 1.

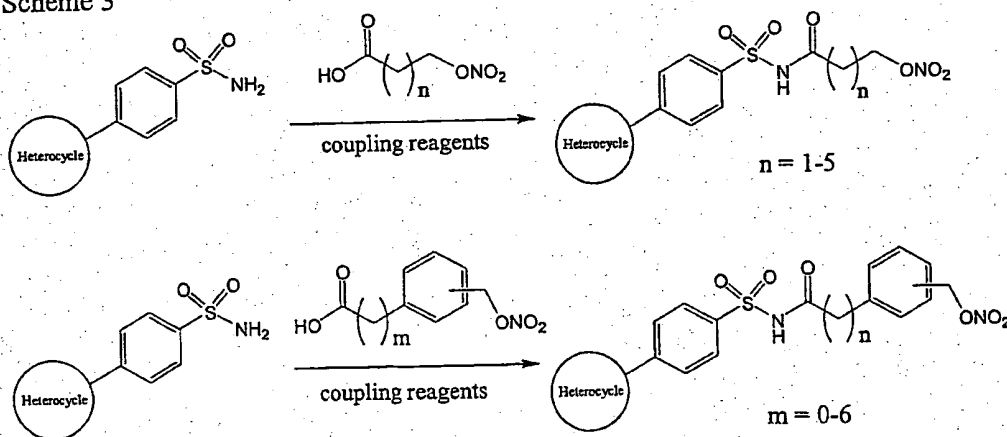
- 10 Reaction of celecoxib or valdecoxib can react with a suitable chlorocarbonate and a base to give the desired product. Alternatively, the desired sulfonylcarbamates can be prepared by converting celecoxib or valdecoxib to bromide derivatives and then nitration of the bromides with silver nitrate (Scheme 2).

Scheme 2



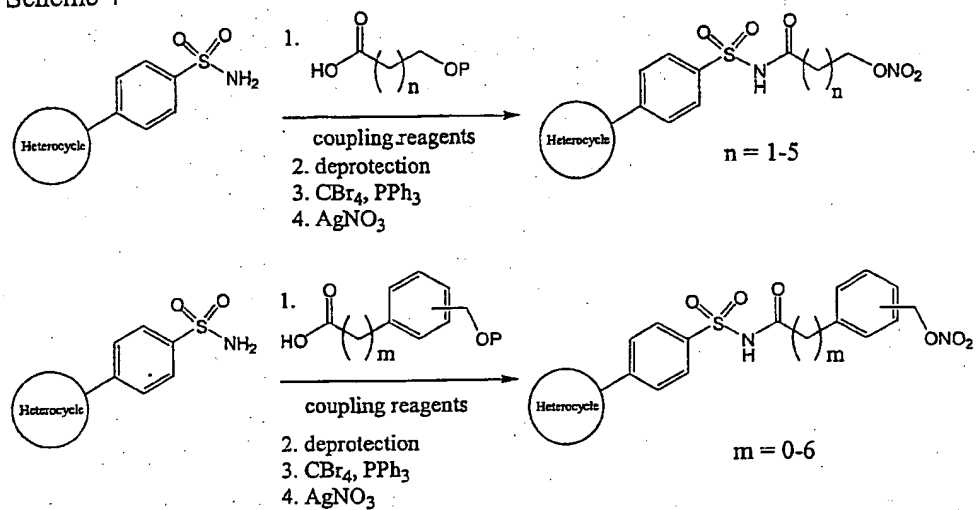
5 Method B

Scheme 3



- The acylsulfonamide derivatives can be prepared by reaction of celecoxib or valdecoxib with the appropriate nitrate containing carboxylic acids (Scheme 3) and a coupling reagent. The standard coupling reagents such as DCC, ECDI, CMC, carbonyldiimidazole or oxalyl chloride can be used for this reaction. Alternatively, celecoxib or valdecoxib can be coupled with the appropriately protected hydroxy-acid derivatives. The resulting intermediates can be deprotected and converted to the corresponding bromides, followed by nitration with silver nitrate to give the desired products (Scheme 4).

Scheme 4



Assays for Determining Biological Activity

The compound of Formula I can be tested using the following assays to determine their biological activity.

5 Inhibition of Cyclooxygenase Activity

Compounds are tested as inhibitors of cyclooxygenase activity in whole cell and microsomal cyclooxygenase assays. Both of these assays measure prostaglandin E₂ (PGE₂) synthesis in response to arachidonic acid, using a radioimmunoassay. Cells used for whole cell assays, and from which microsomes are prepared for microsomal assays, are human
10 osteosarcoma 143 cells (which specifically express cyclooxygenase-2) and human U-937 cells (which specifically express cyclooxygenase-1). In these assays, 100% activity is defined as the difference between prostaglandin E₂ synthesis in the absence and presence of arachidonate addition. IC₅₀ values represent the concentration of putative inhibitor required to return PGE₂ synthesis to 50% of that obtained as compared to the uninhibited control.

15

Representative Rat Paw Edema Assay – Protocol

Male Sprague-Dawley rats (150-200 g) are fasted overnight and are given p.o., either vehicle (1% methocell) or a test compound in the morning. One hr later, a line is drawn using a permanent marker at the level above the ankle in one hind paw to define the area of the
20 paw to be monitored. The paw volume (V_{0h}) is measured using a plethysmometer (Ugo-Basile, Italy) based on the principle of water displacement. The animals are then injected subplantarily with 50 ul of a 1% carrageenan solution in saline (Sigma Chem) into the paw using an insulin syringe with a 25-gauge needle (i.e. 500 ug carrageenan per paw). Three hr later, the paw volume (V_{3h}) is measured and the increases in paw volume (V_{3h} - V_{0h}) are calculated. Paw
25 edema data are compared with the vehicle-control group and percent inhibition calculated taking the values in the control group as 100%. All treatment groups are coded to eliminate observer bias.

NSAID-Induced Gastropathy In Rats

30 Rationale

The major side effect of conventional NSAIDs is their ability to produce gastric lesions in man. Rats are sensitive to the actions of NSAIDs and have been used commonly in the past to evaluate the gastrointestinal side effects of current conventional NSAIDs. In the present assay, NSAID-induced gastrointestinal damage is observed by measuring urinary ⁵¹Cr

excretion after oral dosing of ^{51}Cr -EDTA. Urinary ^{51}Cr excretion is a well-established and sensitive technique to detect gastrointestinal integrity in animals and man.

Methods

- 5 Male Sprague-Dawley rats (150-200 g) are administered orally a test compound either once (acute dosing) or in multiple doses for a few days (chronic dosing). Immediately after the administration of the last dose, the rats are given an oral dose of ^{51}Cr -EDTA (10 $\mu\text{Ci}/\text{rat}$). The animals are placed individually in metabolism cages with food and water *ad lib*. Urine is collected for a 24 hr period and ^{51}Cr urinary excretion is calculated as a percent of
10 total ingested dose.

Protein-Losing Gastropathy in Squirrel Monkeys

Rationale

- 15 Protein-losing gastropathy (manifested as appearance of circulating cells and plasma proteins in the GI tract) is a significant and dose-limiting adverse response to NSAIDs. This can be quantitatively assessed by intravenous administration or $^{51}\text{CrCl}_3$ solution. This isotopic ion can avidly bind to cell and serum globins and cell endoplasmic reticulum. Measurement of radioactivity appearing in feces collected for 24 hr after administration of the
20 isotope thus provides a sensitive and quantitative index of protein-losing gastropathy.

Methods

- Groups of male squirrel monkeys (0.8 to 1.4 kg) are treated by gavage with 1% methocel
25 or a test compounds at multiple doses for a few days. Intravenous ^{51}Cr (5 $\mu\text{Ci}/\text{kg}$ in 1 ml/kg PBS) is administered 1 hr after the last drug/vehicle dose, and feces collected for 24 hr in a metabolism cage and assessed for excreted ^{51}Cr by gamma-counting. ^{51}Cr fecal excretion is calculated as a percent of total injected dose.

Rat Aortic Smooth Muscle Rings in Male Sprague-Dawley Rats

Preparation of rat aortic smooth muscle rings Male Sprague-Dawley rats (Charles River Laboratories (Wilmington, MA) were euthanized by intraperiton injection of a high dose of sodium pentobarbitone (80-100 mg/kg). The thoracic aorta was rapidly excised and immediately placed in a Petri dish containing warm (37 °C) oxygenated (95% O₂ and 5% CO₂) Krebs's buffer (composition per millimolar: NaCl (119); KCl (4.69); CaCl₂·H₂O (2.52); MgSO₄·7H₂O (0.57); NaHCO₃ (25); NaH₂PO₄·H₂O (1.01) and glucose (11.1). Under a stereoscopic dissecting microscope, the aorta was cleaned, freed from adhering fat and connective tissues. The tissue was cut into ring segments, each approximately 2-3 mm in length.

For experiments to measure relaxation of the tissue under various conditions, a stainless steel tissue holder and an U-shaped stainless steel wire were inserted into the lumen of the aortic ring. The tissue holder anchored the ring at the bottom of the organ bath whereas the end of the U-shaped steel wire was tied with fine silk thread so that it connected to the FT-202 transducer. The tissue holder and the steel wire along with the aortic ring were then suspended in a 5-ml, double-jacketed temperature-controlled glass organ bath (Radnoti Glass Technology, Inc., Monrovia, CA) filled with fresh Krebs's buffer. A mixture of 95% O₂ and 5% CO₂ was bubbled through a porous sintered disc at the bottom of the bath. The rings were given an initial resting tension of 1.5 g and the preparation was allowed to equilibrate at the initial tension for about 90 minutes. During this equilibration period, the bath fluid was changed every 15 minutes and replaced with fresh prewarmed (37°C) Krebs's buffer. The isometric tension of the aortic muscle at rest and its response to different stimuli were recorded on a Power Macintosh 6100 computer via a MacLab 8/S computer interface (CB Sciences, Inc, Milford, MA) after an initial amplification through a low-noise ETH-400 bioamplifier (CB Sciences, Inc, Milford, MA). Contractile responsiveness of the tissue strips was established with 10 µM phenylephrine, and the strips were incubated with the drug for 20 minutes to establish a steady level of contraction.

To test the relaxation effects, test compounds were added to the phenylephrine precontracted strips in the tissue bath at cumulative concentrations of 0.1 µM to 0.1 mM. Concentration of test compounds was increased only after relaxation at the previous concentration had reached a plateau level.

Representative Examples

The invention will now be illustrated by the following non-limiting examples in which, unless stated otherwise:

- 5 (i) all operations were carried out at room or ambient temperature, that is, at a temperature in the range 18-25°C; evaporation of solvent was carried out using a rotary evaporator under reduced pressure (600-4000 pascals: 4.5-30 mm. Hg) with a bath temperature of up to 60°C; the course of reactions was followed by thin layer chromatography (TLC) and reaction times are given for illustration
10 only; melting points are uncorrected and 'd' indicates decomposition; the melting points given are those obtained for the materials prepared as described; polymorphism may result in isolation of materials with different melting points in some preparations; the structure and purity of all final products were assured by at least one of the following techniques: TLC, mass spectrometry, nuclear
15 magnetic resonance (NMR) spectrometry or microanalytical data; yields are given for illustration only; when given, NMR data is in the form of delta (δ) values for major diagnostic protons, given in parts per million (ppm) relative to tetramethylsilane (TMS) as internal standard, determined at 300 MHz or 400 MHz using the indicated solvent; conventional abbreviations used for signal
20 shape are: s. singlet; d. doublet; t. triplet; m. multiplet; br. broad; etc.: in addition "Ar" signifies an aromatic signal; chemical symbols have their usual meanings; the following abbreviations have also been used v (volume), w (weight), b.p. (boiling point), m.p. (melting point), L (liter(s)), mL (milliliters), g (gram(s)), mg (milligrams(s)), mol (moles), mmol (millimoles), eq
25 (equivalent(s)).

The following abbreviations have the indicated meanings:

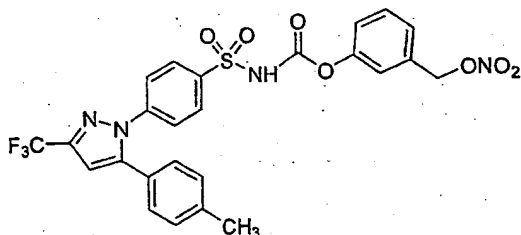
Ac	=	Acetyl
Bn	=	Benzyl
DBU	=	1,8-diazabicyclo[5.4.0]undec-7-ene
DIBAL	=	diisobutylaluminum hydride
DMAP	=	4-(dimethylamino)pyridine
DMF	=	N,N-dimethylformamide

Et ₃ N	=	Triethylamine
HBSS	=	Hanks' balanced salt solution
LDA	=	lithium diisopropylamide
m-CPBA	=	Metachloroperbenzoic acid
MMPP	=	monoperoxyphthalic acid
MPPM	=	monoperoxyphthalic acid, magnesium salt 6H ₂ O
Ms	=	methanesulfonyl = mesyl = S(O) ₂ Me
MsO	=	methanesulfonate = mesylate
NSAID	=	non-steroidal anti-inflammatory drug
OXONE®	=	2KHSO ₅ •KHSO ₄ •K ₂ SO ₄
PBS	=	phosphate buffered saline
PCC	=	pyridinium chlorochromate
PDC	=	pyridinium dichromate
Ph	=	Phenyl
Phe	=	Benzenediyl
Pye	=	Pyridinediyl
r.t.	=	room temperature
rac.	=	Racemic
SAM	=	aminosulfonyl or sulfonamide or S(O) ₂ NH ₂
TBAF	=	tetra-n-butylammonium fluoride
Th	=	2- or 3-thienyl
TFAA	=	trifluoroacetic acid anhydride
THF	=	Tetrahydrofuran
Thi	=	Thiophenediyl
TLC	=	thin layer chromatography
TMS-CN	=	trimethylsilyl cyanide
Tz	=	1H (or 2H)-tetrazol-5-yl
C ₃ H ₅	=	Allyl

Alkyl Group Abbreviations

Me	=	Methyl
Et	=	Ethyl
n-Pr	=	normal propyl
i-Pr	=	Isopropyl
n-Bu	=	normal butyl
i-Bu	=	Isobutyl
s-Bu	=	secondary butyl
t-Bu	=	tertiary butyl
c-Pr	=	Cyclopropyl
c-Bu	=	Cyclobutyl
c-Pen	=	Cyclopentyl
c-Hex	=	Cyclohexyl

EXAMPLE 1

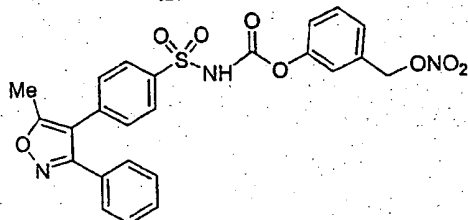


4-[(NITROOXY)METHYL]PHENYL {4-[5-(4-METHYLPHENYL)-3-(TRIFLUOROMETHYL)-1H-PYRAZOL-1-YL]PHENYL} SULFONYLCARBAMATE

To a solution of 3-nitrooxymethylphenol (1.69 g, 10 mmol) and triphosgene (0.92 g, 3.1 mmol) in 100 ml of CH₂Cl₂, pyridine (0.86 g, 11 mmol) is added dropwise at -78°C. After 1 h of stirring at rt, the final suspension is transferred via a canula to a solution of Celebrex (3.81 g, 10 mmol) and pyridine 0.86g in CH₂Cl₂ cooled at -78°C. After 1 h at 0°C, the reaction mixture is poured into a separatory funnel containing EtOAc(50 mL)/ NH₄Cl(50 mL, sat). The phases are separated and the aqueous phase is extracted twice with EtOAc (2 x 50 mL). The organic layers are combined, washed with brine, dried over anhydrous Na₂SO₄ and evaporated to dryness. The

crude material can be further purified by flash chromatography eluting with 1:3 EtOAc / hexanes to yield the title compound.

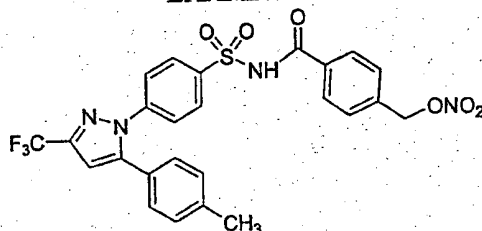
EXAMPLE 2



4-[(NITROOXY)METHYL]PHENYL [4-(5-METHYL-3-PHENYLISOXAZOL-4-YL)PHENYL]SULFONYLCARBAMATE

Starting from valdecoxib, the title compound is prepared by following the same conditions as described in Example 1.

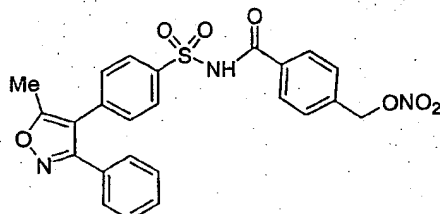
EXAMPLE 3



4-[[[4-[5-(4-METHYLPHENYL)-3-(TRIFLUOROMETHYL)-1H-PYRAZOL-1-YL]PHENYL]SULFONYL]AMINO]CARBONYL}BENZYL NITRATE

To a solution of 4-nitrooxymethylbenzoic acid (1.97 g, 10 mmol) and Celebrex (3.81 g, 10 mmol) in 100 ml of CH_2Cl_2 is added EDCI (3.8 g, 20 mmol) at 0°C . After 1h of stirring at rt, the reaction mixture is poured into a separatory funnel containing EtOAc(50 mL)/ NH_4Cl (50 mL, sat). The phases are separated and the aqueous phase is extracted twice with EtOAc (2 x 50 mL). The organic layers are combined, washed with brine, dried over anhydrous Na_2SO_4 and evaporated to dryness. The crude material can be further purified by flash chromatography eluting with 1:3 EtOAc / hexanes to yield the title compound.

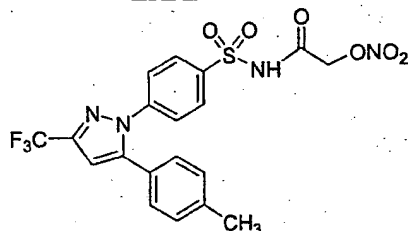
EXAMPLE 4



4-[(4-(5-METHYL-3-PHENYLISOXAZOL-4-
YL)PHENYL]SULFONYL}AMINO)CARBONYL]BENZYL NITRATE

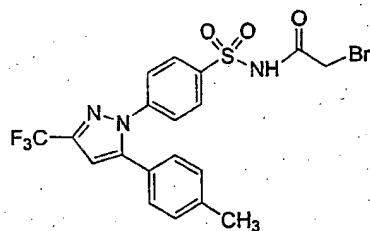
Starting from valdecixib, the title compound is prepared by following the same conditions as described in Example 3.

EXAMPLE 5



2-[(4-[5-(4-METHYLPHENYL)-3-(TRIFLUOROMETHYL)-1H-PYRAZOL-1-
YL]PHENYL}SULFONYL)AMINO]-2-OXOETHYL NITRATE

Step 1

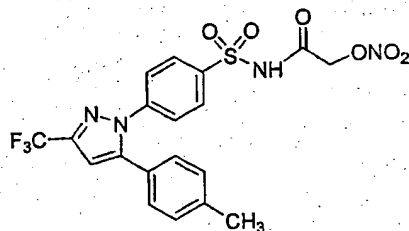


N-(Bromoacetyl)-4-[5-(4-methylphenyl)-3-(trifluoromethyl)-1*H*-pyrazol-1-yl]benzenesulfonamide

To a solution of Celebrex (1.9 g, 5 mmol) and Et₃N (0.7 g, 7 mmol) in 30 ml of CH₂Cl₂ is added 2-bromoacetyl bromide (1.0g, 5 mmol) at 0°C. After 1h of stirring at rt, the reaction mixture is poured into a separatory funnel containing EtOAc(50 mL)/ NH₄Cl(50 mL, sat). The

phases are separated and the aqueous phase is extracted twice with EtOAc (2 x 50 mL). The organic layers are combined, washed with brine, dried over anhydrous Na₂SO₄ and evaporated to dryness. The crude material can be further purified by flash chromatography eluting with 1:3 EtOAc / hexanes to yield the title compound.

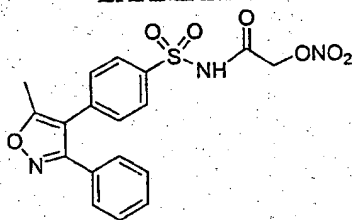
Step 2



2-[(4-{5-(4-Methylphenyl)-3-(trifluoromethyl)-1*h*-pyrazol-1-yl}phenyl)sulfonyl]amino-2-oxoethyl nitrate

To a solution of the product of Step 1 in 40 mL of acetonitrile is added AgNO₃ (1.7 g, 10 mmol) at room temperature. The reaction mixture is stirred for 30 min and then diluted with 100 mL of 1:1 EtOAc/hexane. The resulting suspension is filtered through a pad of silica gel and the filtrate is concentrated and the residue is purified by silica gel chromatography eluted with 2:1 hexane/EtOAc to give the title compound.

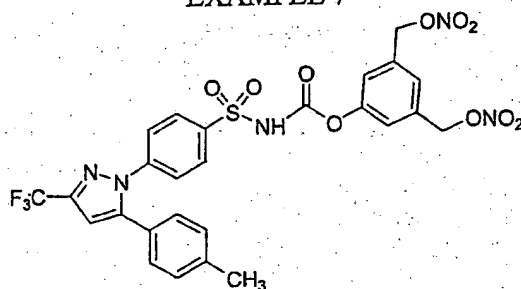
EXAMPLE 6



2-[(4-{5-METHYL-3-PHENYLISOXAZOL-4-YL}PHENYL)SULFONYL]AMINO-2-OXOETHYL NITRATE

Starting from valdecoxib, the title compound is prepared by following the same conditions as described in Example 5.

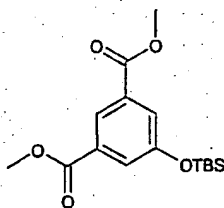
EXAMPLE 7



3,5-BIS[(NITROOXY)METHYL]PHENYL {4-[5-(4-METHYLPHENYL)-3-(TRIFLUOROMETHYL)-1H-PYRAZOL-1-YL]PHENYL} SULFONYLCARBAMATE

5

Step 1



Dimethyl 5-[[tert-butyl(dimethyl)silyl]oxy]isophthalate

10

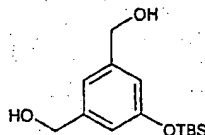
A solution of 14.8 g of dimethyl 5-hydroxyisophthalate, 11.7 g of tert-butyldimethylsilyl chloride and 5.76 g of imidazole in 125 ml of DMF was stirred at 25°C overnight under N₂. To quench the reaction, 300 ml of NH₄Cl sat and 300 ml of ethyl acetate were added and the mixture was stirred 10 min. The phases were separated and the aqueous phase was extracted twice with 100 ml of ethyl acetate. The organic layers were combined, washed with brine, dried over anhydrous Na₂SO₄, filtered and concentrated to yield 21.9 g of the title compound as a white powder. The crude material was used directly for the next step without further purification.

15

¹H NMR (acetone-d₆) δ 8.22 (s, 1H), 7.67 (s, 2H), 3.91 (s, 6H), 0.99 (s, 9H), 0.26 (s, 6H).

20

Step 2

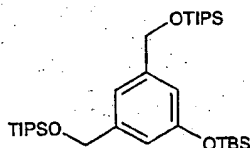


3-(tert-Butyl(dimethyl)silyloxy)-5-hydroxymethylphenyl methanol

To a solution of 10.2 g of the previous compound in 100 mL of anhydrous THF under N₂ at -10°C was added 135 mL of DIBAL-H (1M/hexanes) via a dropping funnel. The reaction was slowly warm-up to room temperature and stirred 3h. To quench the reaction, 500 ml of Rochelle's salt and 300 ml of ethyl acetate were added and the mixture was stirred overnight. The phases were separated and the aqueous phase was extracted twice with 300 ml of ethyl acetate. The organic layers were combined, washed with brine, dried over anhydrous Na₂SO₄, filtered and concentrated to yield 8.37 g of the title compound as a white powder. The crude material was used directly for the next step without further purification.

¹H NMR (acetone-d₆) δ 6.90 (s, 1H), 6.74 (s, 2H), 4.55 (d, 4H), 4.12 (t, 2H), 0.99 (s, 9H), 0.20 (s, 6H).

Step 3

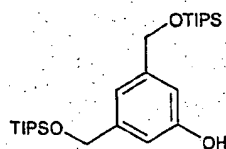


1-(tert-Butyl(dimethyl)silyloxy)-3,5-bis-triisopropylsilyloxymethyl benzene

A solution of 7.0 g of the previous compound, 12.3 mL of triisopropylsilyl chloride and 4.25 g of imidazole in 100 ml of DMF was stirred at 25°C overnight under N₂. To quench the reaction, 300 ml of NH₄Cl sat and 300 ml of ethyl acetate were added and the mixture was stirred 10 min. The phases were separated and the aqueous phase was extracted twice with 100 ml of ethyl acetate. The organic layers were combined, washed with brine, dried over anhydrous Na₂SO₄, filtered and concentrated to yield 13.9 g of the title compound as a colorless oil. The crude material was used directly for the next step without further purification.

¹H NMR (acetone-d₆) δ 7.01 (s, 1H), 6.83 (s, 2H), 4.85 (s, 4H), 1.22 (bs, 6H), 1.12 (d, 36H), 0.99 (s, 9H), 0.22 (s, 6H).

Step 4

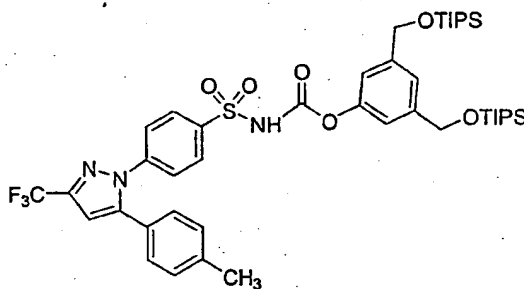


3,5-Bis-triisopropylsilyloxymethyl phenol

To a solution of 12.7 g of the previous compound in 120 ml of THF was added 22.2 mL of TBAF (1M / THF) at 0°C under N₂. The reaction was stirred 15 min then quenched by adding 200 ml of NH₄Cl sat and 300 ml of diethyl ether were added and the mixture was stirred 15 min. The phases were separated and the aqueous phase was extracted twice with 100 ml of diethyl ether. The organic layers were combined, washed with brine, dried over anhydrous Na₂SO₄, filtered and concentrated to yield 9.65 g of the title compound as a colorless oil. The crude material was used directly for the next step without further purification

¹H NMR (acetone-d₆) δ 8.27 (s, 1H), 6.90 (s, 1H), 6.78 (s, 2H), 4.82 (s, 4H), 1.22 (bs, 6H), 1.12 (d, 36H).

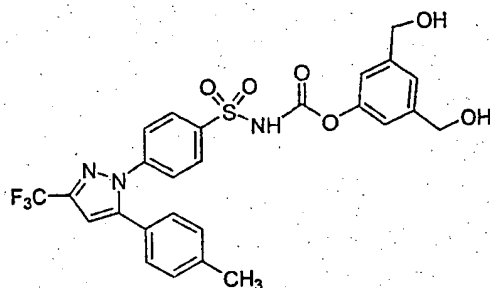
Step 5



3,5-Bis[(triisopropylsilyloxy)methyl]phenyl {4-[5-(4-methylphenyl)-3-(trifluoromethyl)-1H-pyrazol-1-yl]phenyl}sulfonylcarbamate

To a solution of the previous phenol (product of Step 4) and triphosgene in CH₂Cl₂, pyridine is added dropwise at -78°C. After 1h of stirring at rt, the final suspension is transferred via a canula to a solution Celebrex and pyridine mL CH₂Cl₂ cooled at -78°C. After 1 h at 0°C, the reaction mixture is poured into a separatory funnel containing EtOAc/ NH₄Cl(sat). The phases were separated and the aqueous phase is extracted twice with EtOAc. The organic layers are combined, washed with brine, dried over anhydrous Na₂SO₄ and evaporated to dryness. The crude material is further purified by flash chromatography eluting with EtOAc / hexanes to yield the title compound.

Step 6



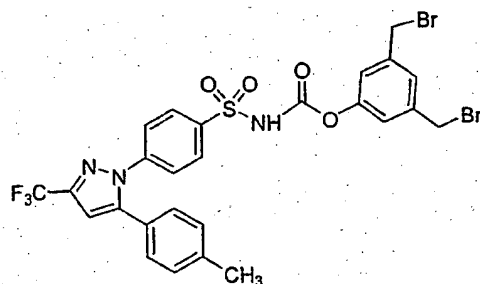
3,5-Bis(hydroxymethyl)phenyl {4-[5-(4-methylphenyl)-3-(trifluoromethyl)-1*H*-pyrazol-1-yl]phenyl} sulfonylcarbamate

5

A solution of the product of Step 5 at rt in acetonitrile under a nitrogen flow is added HF/pyridine complex (70% HF) over 5 min. After 1h at room temperature, the reaction mixture is poured into a separatory funnel containing EtOAc/ CuSO₄(sat). The phases are separated and the aqueous phase is extracted twice with EtOAc. The organic layers are combined, washed with CuSO₄(sat), brine, dried over anhydrous Na₂SO₄ and evaporated to dryness. The crude material is further purified by flash chromatography eluting with EtOAc / hexanes to yield the title compound.

10

Step 7



15

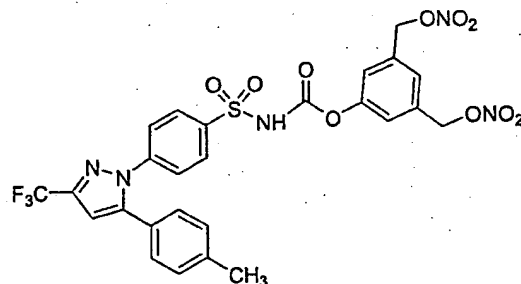
3,5-Bis(bromomethyl)phenyl {4-[5-(4-methylphenyl)-3-(trifluoromethyl)-1*H*-pyrazol-1-yl]phenyl} sulfonylcarbamate

20

To a solution of triphenylphosphine in CH₂Cl₂ is added bromine (1M / CH₂Cl₂) at 0°C. After 10 min, Hünig's base is added followed by the previous alcohol in CH₂Cl₂. After warming to room temperature, the reaction mixture is poured into a separatory funnel containing EtOAc/brine. The phases are separated and the organic layer is dried over anhydrous Na₂SO₄

and evaporated to dryness. The crude material is further purified by flash chromatography eluting with EtOAc / hexanes to yield the title compound.

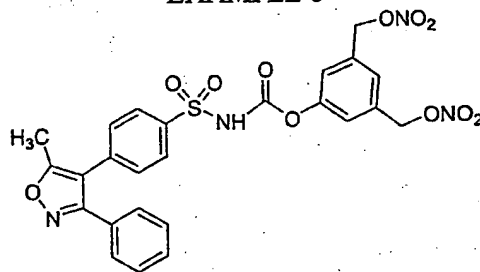
Step 8



3,5-Bis[(nitrooxy)methyl]phenyl {4-[5-(4-methylphenyl)-3-(trifluoromethyl)-1*h*-pyrazol-1-yl]phenyl} sulfonylcarbamate

To a solution of the product of Step 7 in acetonitrile silver nitrate is added. After stirring for 2 h at room temperature the reaction mixture is diluted with 1:1 EtOAc/hexane and filtered through a pad of silica gel eluting with EtOAc. The solvents were removed under vacuo. The crude material was further purified by flash chromatography eluting with EtOAc / hexanes to yield the title compound.

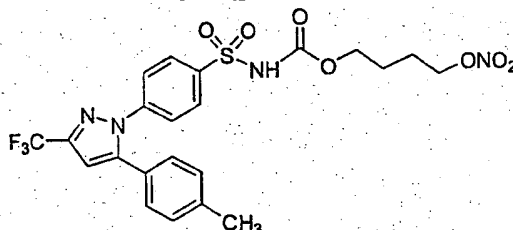
EXAMPLE 8



3,5-BIS[(NITROOXY)METHYL]PHENYL [4-(5-METHYL-3-PHENYLISOXAZOL-4-YL)PHENYL]SULFONYLCARBAMATE

Starting from valdecoxib, the title compound is prepared by following the same conditions as described in Step 5-8 of Example 7.

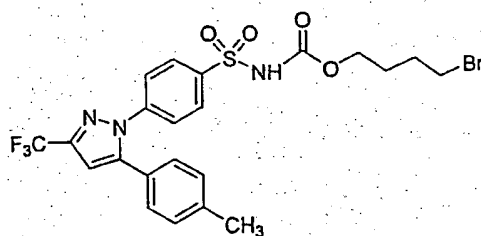
EXAMPLE 9



4-(NITROOXY)BUTYL {4-[5-(4-METHYLPHENYL)-3-(TRIFLUOROMETHYL)-1H-PYRAZOL-1-YL]PHENYL} SULFONYLCARBAMATE

5

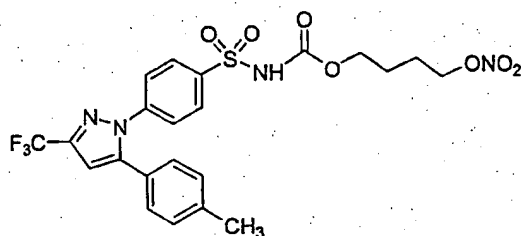
Step 1



10 4-Bromobutyl {4-[5-(4-methylphenyl)-3-(trifluoromethyl)-1H-pyrazol-1-yl]phenyl} sulfonycarbamate

To a solution of 4-bromobutanol (1.53 g, 10 mmol) and triphosgene (0.91 g, 3 mmol) in 100 ml of CH₂Cl₂, pyridine (0.86 g, 11 mmol) is added dropwise at -78°C. After 1h of stirring at
 15 rt, the final suspension is transferred via a canula to a solution of Celebrex (3.81 g, 10 mmol) and pyridine 0.86g in CH₂Cl₂ cooled at -78°C. After 15 min at 0°C, the reaction mixture is poured into a separatory funnel containing EtOAc(50 mL)/ NH₄Cl(50 mL, sat). The phases are separated and the aqueous phase is extracted twice with EtOAc (2 x 50 mL). The organic layers are combined, washed with brine, dried over anhydrous Na₂SO₄ and evaporated to dryness. The
 20 crude material can be further purified by flash chromatography eluting with 1:3 EtOAc / hexanes to yield the title compound.

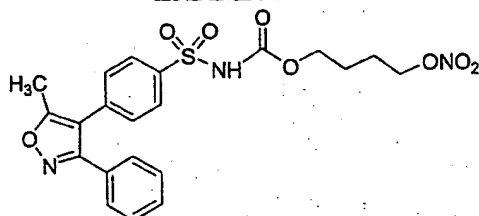
Step 2



5 4-(Nitrooxy)butyl {4-[5-(4-methylphenyl)-3-(trifluoromethyl)-1*h*-pyrazol-1-yl]phenyl} sulfonylcarbamate

To a solution of the product of Step 1 in 40 mL of acetonitrile is added AgNO₃ (1.9 g, 20 mmol) at room temperature. The reaction mixture is stirred for 30 min and then diluted with 100 mL of 1:1 EtOAc/hexane. The resulting suspension is filtered through a pad of silica gel and the
 10 filtrate is concentrated and the residue is purified by silica gel chromatography eluted with 2:1 hexane/EtOAc to give the title compound.

EXAMPLE 10



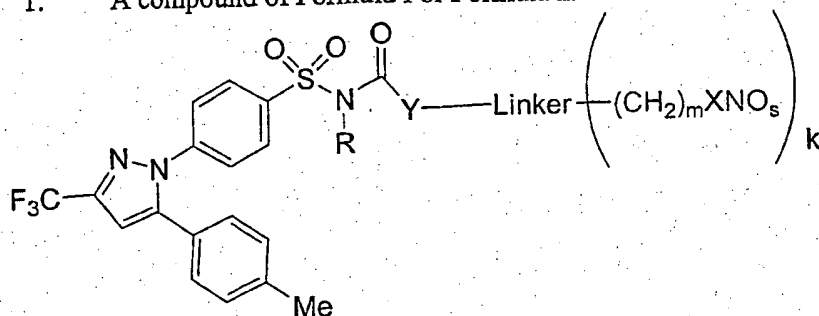
15 4-(NITROOXY)BUTYL [4-(5-METHYL-3-PHENYLISOXAZOL-4-YL)PHENYL]SULFONYLCARBAMATE

Starting from valdecoxib, the title compound is prepared by following the same conditions as described in Step 1-2 of Example 9.

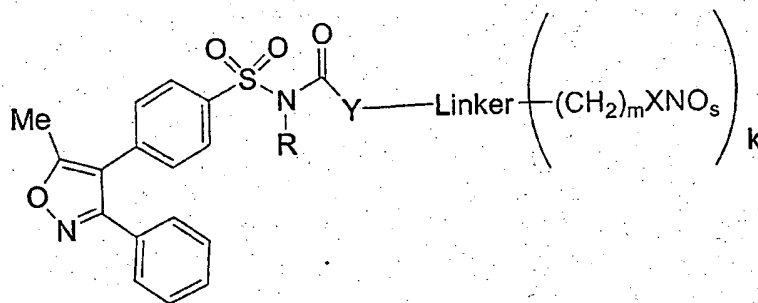
20

WHAT IS CLAIMED IS:

1. A compound of Formula I or Formula II



Formula I



Formula II

or a pharmaceutically acceptable salt thereof wherein

each s is independently 1 or 2;

k is 1, 2, 3 or 4;

each m is independently 0, 1, 2, 3 or 4;

each X is independently O or S;

Y is a bond, S, O or NR₁, wherein R₁ is hydrogen or C₁-6alkyl;

R is hydrogen or C₁-6alkyl;

the Linker is selected from the group consisting of:

- (a) $-(CH_2)_n$, wherein n is 0, 1, 2, 3 or 4,

- (b) C₃₋₆cycloalkyl, wherein the C₃₋₆cycloalkyl optionally mono-, di- or tri-substituted with a substituent selected from the group consisting of (1) halo,
(2) C₁₋₃alkyl,
(3) C₁₋₃alkoxy,
5 (4) Hydroxy,
(5) NO₂,
(6) CO₂,
(7) CF₃,
(8) CN;
10 (9) CH₂COOH
(10) CH₂COO-C₁₋₃alkyl,
(11) C₁₋₃alkthio,
- (c) aryl, wherein the aryl is selected from the group consisting of phenyl and naphthyl, wherein the aryl is optionally mono-, di- or tri-substituted with a
15 substituent selected from the group consisting of
(1) halo,
(2) C₁₋₃alkyl,
(3) C₁₋₃alkoxy,
(4) Hydroxy,
20 (5) NO₂,
(6) CO₂,
(7) CF₃,
(8) CN;
(9) CH₂COOH
25 (10) CH₂COO-C₁₋₃alkyl,
(11) C₁₋₃alkthio,
- (d) Heteroaryl optionally mono-, di- or tri- substituted with substituents selected
from the group consisting of,
(1) halo,
30 (2) C₁₋₃alkyl,
(3) C₁₋₃alkoxy,
(4) Hydroxy,
(5) NO₂,
(6) CO₂,

- (7) CF₃,
(8) CN;
(9) CH₂COOH
(10) CH₂COO-C₁₋₃alkyl,
5 (11) C₁₋₃alkthio.

2. The compound according to Claim 1 wherein

s is 2;

k is 1;

10 m is 1 or 2.

3. The compound according to Claim 1 wherein

X is O;

R is H.

15

4. The compound according to Claim 1 wherein

R is H; and

Y is a bond.

20

5. The compound according to Claim 1 wherein

R is H;

Y is a bond;

s is 2;

k is 1;

25 m is 1.

6. The compound according to Claim 2 wherein:
the Linker is -(CH₂)_n, wherein n is 1 or 2.

30

7. The compound according to Claim 2 wherein:
the Linker is C₃₋₆cycloalkyl, wherein the C₃₋₆cycloalkyl optionally mono-, di- or tri-substituted
with a substituent selected from the group consisting of

(1) halo,

(2) Methyl,

- (3) Methoxy,
- (4) Hydroxy,
- (5) NO₂,
- (6) CO₂,
- (7) CF₃,
- (8) CN, and
- (9) CH₂COOH.

5

8. The compound according to Claim 2 wherein:

10

the Linker is aryl, wherein the aryl is selected from the group consisting of phenyl and naphthyl, wherein the aryl is optionally mono-, di- or tri-substituted with a substituent selected from the group consisting of

- (1) halo,
- (2) Methyl,
- (3) Methoxy,
- (4) Hydroxy,
- (5) NO₂,
- (6) CO₂,
- (7) CF₃,
- (8) CN, and
- (9) CH₂COOH.

15

20

9. The compound according to Claim 2 wherein:

25

the Linker is phenyl optionally mono-, di- or tri-substituted with a substituent selected from the group consisting of:

- (1) halo,
- (2) Methyl,
- (3) Methoxy,
- (4) Hydroxy,
- (5) NO₂,
- (6) CO₂,
- (7) CF₃,
- (8) CN, and
- (9) CH₂COOH.

30

10. The compound according to Claim 2 wherein:

the Linker is benzimidazolyl, benzofuranyl, benzopyrazolyl, benzotriazolyl,
benzothiophenyl, benzoxazolyl, carbazolyl, carbolinyl, cinnolyl, furanyl,
imidazolyl, indolyl, indolyl, indolaziny, indazolyl, isobenzofuranyl, isoindolyl,
isoquinolyl, isothiazolyl, isoxazolyl, naphthyridinyl, oxadiazolyl, oxazolyl,
pyrazinyl, pyrazolyl, pyridopyridinyl, pyridazinyl, pyridyl, pyrimidyl, pyrrolyl,
quinazolinyl, quinolyl, quinoxalyl, thiadiazolyl, thiazolyl, thienyl, triazolyl,
azetidyl, 1,4-dioxanyl, hexahydroazepinyl, piperazinyl, piperidinyl,
pyrrolidinyl, morpholinyl, thiomorpholinyl, dihydrobenzimidazolyl,
dihydrobenzofuranyl, dihydrobenzothiophenyl, dihydrobenzoxazolyl,
dihydrofuranyl, dihydroimidazolyl, dihydroindolyl, dihydroisooxazolyl,
dihydroisothiazolyl, dihydrooxadiazolyl, dihydrooxazolyl, dihydropyrazinyl,
dihydropyrazolyl, dihydropyridinyl, dihydropyrimidinyl, dihydropyrrolyl,
dihydroquinolyl, dihydrotetrazolyl, dihydrothiadiazolyl, dihydrothiazolyl,
dihydrothienyl, dihydrotriazolyl, dihydroazetidyl, methylenedioxybenzoyl,
tetrahydrofuranyl, and tetrahydrothienyl, optionally mono-, di- or tri- substituted
with substituents selected from the group consisting of,

- (1) halo,
- (2) Methyl,
- (3) Methoxy,
- (4) Hydroxy,
- (5) NO₂,
- (6) CO₂,
- (7) CF₃,
- (8) CN, and
- (9) CH₂COOH.

11. The compound according to Claim 10 wherein the Linker is selected from
pyridyl, each optionally mono-, di- or tri- substituted with substituents selected
from the group consisting of,

- (1) halo,
- (2) Methyl,

- (3) Methoxy,
(4) Hydroxy,
(5) NO₂,
(6) CO₂,
5 (7) CF₃,
(8) CN, and
(9) CH₂COOH.

12. The compound according to Claim 1 wherein s is 2.

10 13. A method of treating an inflammatory disease susceptible to treatment with a non-steroidal anti-inflammatory agent comprising administering to a patient in need of such treatment of a non-toxic therapeutically effective amount of a compound according to Claim 1.

15 14. The method according to Claim 13 wherein the patient is also at risk of a thrombotic cardiovascular event.

20 15. A method of treating cyclooxygenase mediated diseases advantageously treated by an active agent that selectively inhibits COX-2 in preference to COX-1 comprising administering to a patient in need of such treatment of a non-toxic therapeutically effective amount of a compound according to Claim 1.

25 16. The method according to Claim 15 wherein the patient is also at risk of a thrombotic cardiovascular event.

30 17. A method for treating a chronic cyclooxygenase-2 mediated disease or condition and reducing the risk of a thrombotic cardiovascular event in a human patient in need of such treatment and at risk of a thrombotic cardiovascular event comprising orally concomitantly or sequentially administering to said patient a compound according to Claim 1 in an amount effective to treat the cyclooxygenase-2 mediated disease or condition and aspirin in an amount effective to reduce the risk of the thrombotic cardiovascular event.

18. The method according to Claim 17 wherein the compound is administered orally on a once daily basis.

5 19. The method according to Claim 17 wherein the compound is administered orally on a twice daily basis.

10 20. The method according to Claim 17 wherein the cyclooxygenase-2 selective mediated disease or condition is selected from the group consisting of: osteoarthritis, rheumatoid arthritis and chronic pain.

21. The method according to Claim 17 wherein aspirin is administered at a dose of about 30 mg to about 1 g.

15 22. The method according to Claim 17 wherein aspirin is administered at a dose of about 80 to about 650 mg.

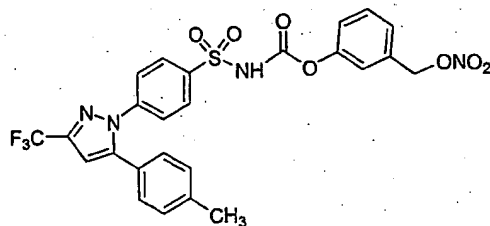
23. The method according to Claim 22 wherein aspirin is administered at a dose of about 81 mg or about 325 mg.

20 24. The method according to Claim 22 wherein aspirin is orally administered once daily.

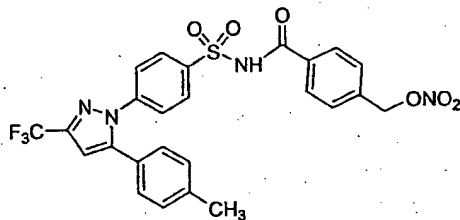
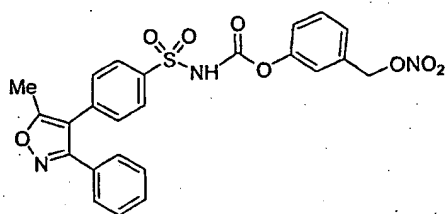
25 25. A pharmaceutical composition comprising a compound according to any one of Claims 1 to 12 and aspirin in combination with a pharmaceutically acceptable carrier.

26. A pharmaceutical composition comprising a compound according to any one of Claims 1 to 12 and a pharmaceutically acceptable carrier.

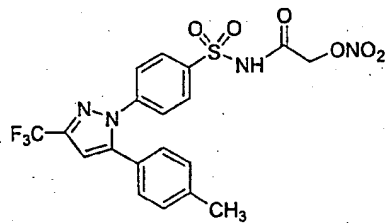
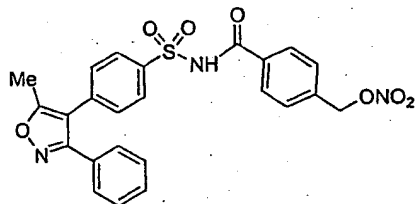
27. A compound selected from the following group:

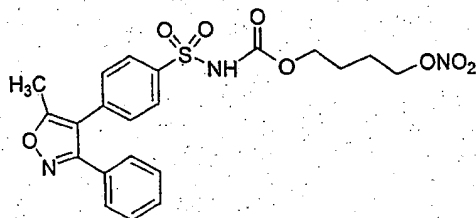
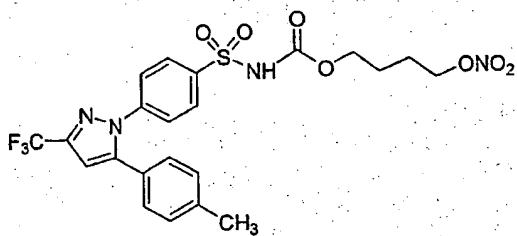
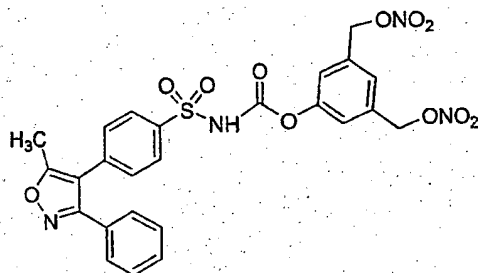
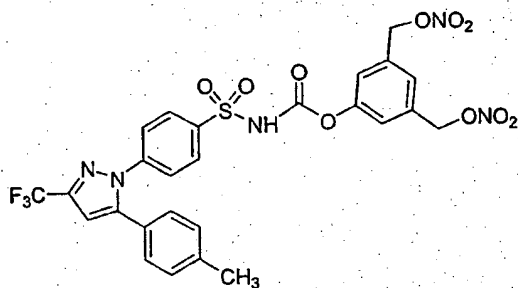
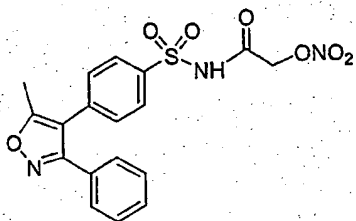


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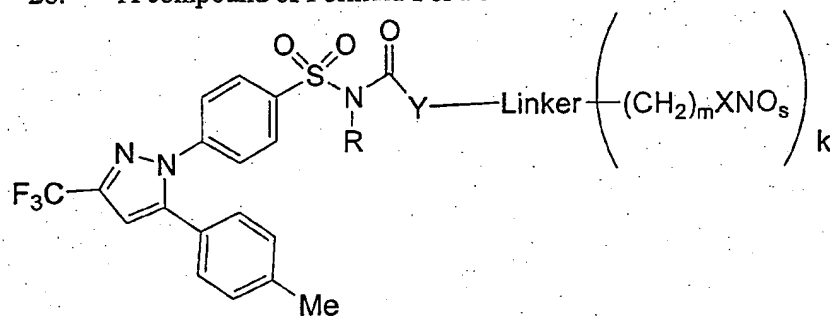


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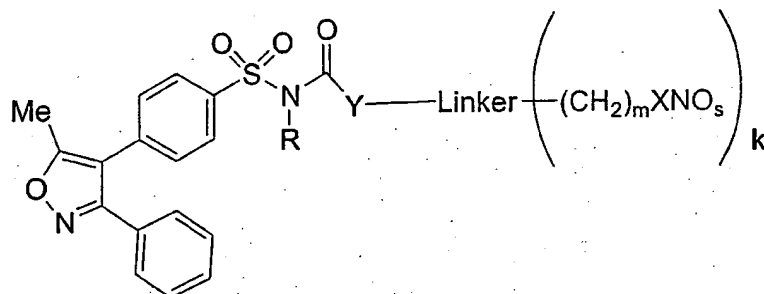




28. A compound of Formula I or Formula II



Formula I



Formula II

or a pharmaceutically acceptable salt thereof wherein
each s is independently 1 or 2;

10 k is 1, 2, 3 or 4;

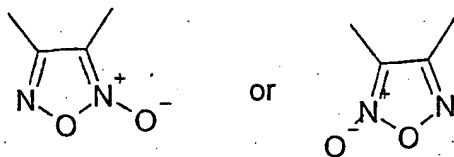
each m is independently 0, 1, 2, 3 or 4;

each X is independently O or S;

Y is a bond, S, O or NR₁, wherein R₁ is hydrogen or C₁-6alkyl;

R is hydrogen or C₁-6alkyl;

15 the Linker is selected from the group consisting of:



29. Use of a compound of any one of Claims 1 to 12, 27 or 28, or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for treating an inflammatory disease.

30. A compound of any one of Claims 1 to 12, 27 or 28, or a pharmaceutically acceptable salt thereof, for use in medicinal therapy.

31. A compound of any one of Claims 1 to 12, 27 or 28, or a pharmaceutically acceptable salt thereof, for use in treating cyclooxygenase mediated diseases advantageously treated by an agent that selectively inhibits COX-2 in preference to COX-1.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/CA 03/01605

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D261/08 C07D231/12 A61K31/415 A61K31/42

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D A61K A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, CHEM ABS Data, WPI Data, EMBASE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 01/045703 A (FANG XINQIN ; GARVEY DAVID S (US); LETTS L GORDON (US); NITROMED INC () 28 June 2001 (2001-06-28) page 4, line 25 - page 8, line 10; claim 1; example 16	1-31
Y	WO 00/25776 A (DONG QING ; FANG XINQIN (US); GARVEY DAVID S (US); NITROMED INC (US);) 11 May 2000 (2000-05-11) page 2, line 26 - page 5, line 14; claim 1	1-31
Y	US 5 633 272 A (STEALEY MICHAEL A ET AL) 27 May 1997 (1997-05-27) example 1	1-31
Y	US 5 466 823 A (GRANETO MATTHEW J ET AL) 14 November 1995 (1995-11-14) example 1c	1-31
	-/-	



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- *Z* document member of the same patent family

Date of the actual completion of the international search

9 February 2004

Date of mailing of the international search report

16/02/2004

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Authorized officer

Usueli, A

INTERNATIONAL SEARCH REPORT

International Application No

PCT/CA 03/01605

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	EP 1 336 602 A (SCARAMUZZINO GIOVANNI) 20 August 2003 (2003-08-20) see compounds E1a, E1b, E1d, EVa, EVd, EVeclaim 1 page 83, column 84 -----	1-31
E	WO 2004/000781 A (NICOX SA ; DEL SOLDATO PIERO (IT); SANTUS GIANCARLO (IT)) 31 December 2003 (2003-12-31) claim 1; example 4 -----	1-31

INTERNATIONAL SEARCH REPORT

International application No.
PCT/CA 03/01605

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
Although claims 13-24 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/CA 03/01605

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 0145703	A	28-06-2001	AU 2592801 A	03-07-2001
			BR 0017037 A	10-06-2003
			CA 2393724 A1	28-06-2001
			CN 1434712 T	06-08-2003
			EP 1246621 A1	09-10-2002
			JP 2003523958 T	12-08-2003
			WO 0145703 A1	28-06-2001
			US 2003220228 A1	27-11-2003
US 2001041726 A1	15-11-2001			
WO 0025776	A	11-05-2000	AU 763000 B2	10-07-2003
			AU 1601200 A	22-05-2000
			CA 2348741 A1	11-05-2000
			EP 1126838 A1	29-08-2001
			JP 2002528495 T	03-09-2002
			WO 0025776 A1	11-05-2000
			US 6297260 B1	02-10-2001
			US 2003207919 A1	06-11-2003
US 2002016322 A1	07-02-2002			
US 5633272	A	27-05-1997	AT 223390 T	15-09-2002
			AU 699593 B2	10-12-1998
			AU 4867196 A	04-09-1996
			BR 9607035 A	04-11-1997
			CA 2212836 A1	22-08-1996
			CN 1442139 A	17-09-2003
			CN 1181075 A ,B	06-05-1998
			CZ 9702546 A3	14-01-1998
			DE 69623444 D1	10-10-2002
			DE 69623444 T2	05-06-2003
			DK 809636 T3	06-01-2003
			EP 1223167 A2	17-07-2002
			EP 0809636 A1	03-12-1997
			ES 2183935 T3	01-04-2003
			FI 973292 A	10-10-1997
			JP 3267300 B2	18-03-2002
			JP 11503722 T	30-03-1999
			JP 2002179656 A	26-06-2002
			LU 91024 A9	04-08-2003
			NO 973711 A	06-10-1997
			NZ 302586 A	30-08-1999
			PL 321814 A1	22-12-1997
			PL 185544 B1	30-05-2003
			PT 809636 T	31-12-2002
			RU 2200158 C2	10-03-2003
			WO 9625405 A1	22-08-1996
			US 5859257 A	12-01-1999
			US 5985902 A	16-11-1999
ZA 9601150 A	12-02-1997			
US 5466823	A	14-11-1995	AT 187965 T	15-01-2000
			AT 212985 T	15-02-2002
			AT 233245 T	15-03-2003
			AT 219937 T	15-07-2002
			AU 690609 B2	30-04-1998
			AU 1171495 A	19-06-1995
			BR 1100406 A3	08-02-2000
			CA 2177576 A1	08-06-1995

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/CA 03/01605

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
US 5466823	A		CN 1141630 A ,B	29-01-1997
			CN 1280125 A ,B	17-01-2001
			CN 1280126 A ,B	17-01-2001
			CY 2237 A	04-07-2003
			CZ 9601503 A3	11-12-1996
			DE 69422306 D1	27-01-2000
			DE 69422306 T2	18-05-2000
			DE 69429836 D1	21-03-2002
			DE 69429836 T2	18-07-2002
			DE 69430930 D1	08-08-2002
			DE 69430930 T2	20-02-2003
			DE 69432193 D1	03-04-2003
			DE 69432193 T2	15-01-2004
			DK 731795 T3	15-05-2000
			DK 924201 T3	21-05-2002
			DK 922697 T3	10-06-2003
			DK 923933 T3	21-10-2002
			EP 0731795 A1	18-09-1996
			EP 0924201 A1	23-06-1999
			EP 0922697 A1	16-06-1999
			EP 0923933 A1	23-06-1999
			ES 2141916 T3	01-04-2000
			ES 2172959 T3	01-10-2002
			ES 2193609 T3	01-11-2003
			ES 2180233 T3	01-02-2003
			FI 962249 A	29-05-1996
			GR 3032696 T3	30-06-2000
			HK 1013649 A1	07-07-2000
			HK 1021935 A1	04-04-2003
			HU 74180 A2	28-11-1996
			JP 3445762 B2	08-09-2003
			JP 2000109466 A	18-04-2000
			JP 3025017 B2	27-03-2000
			JP 9506350 T	24-06-1997
			JP 2003238536 A	27-08-2003
			KR 229343 B1	01-11-1999
			KR 263817 B1	16-08-2000
			KR 261669 B1	15-07-2000
			LU 90698 A9	13-02-2001
			NO 962184 A	29-05-1996
			NZ 276885 A	30-08-1999
			PL 314695 A1	16-09-1996
EP 1336602	A	20-08-2003	EP 1336602 A1	20-08-2003
WO 2004000781	A	31-12-2003	IT MI20021391 A1	29-12-2003
			WO 2004000781 A2	31-12-2003

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